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(54) **METHOD AND APPARATUS FOR CONTROLLING THE DEPLOYMENT OF A STENT**

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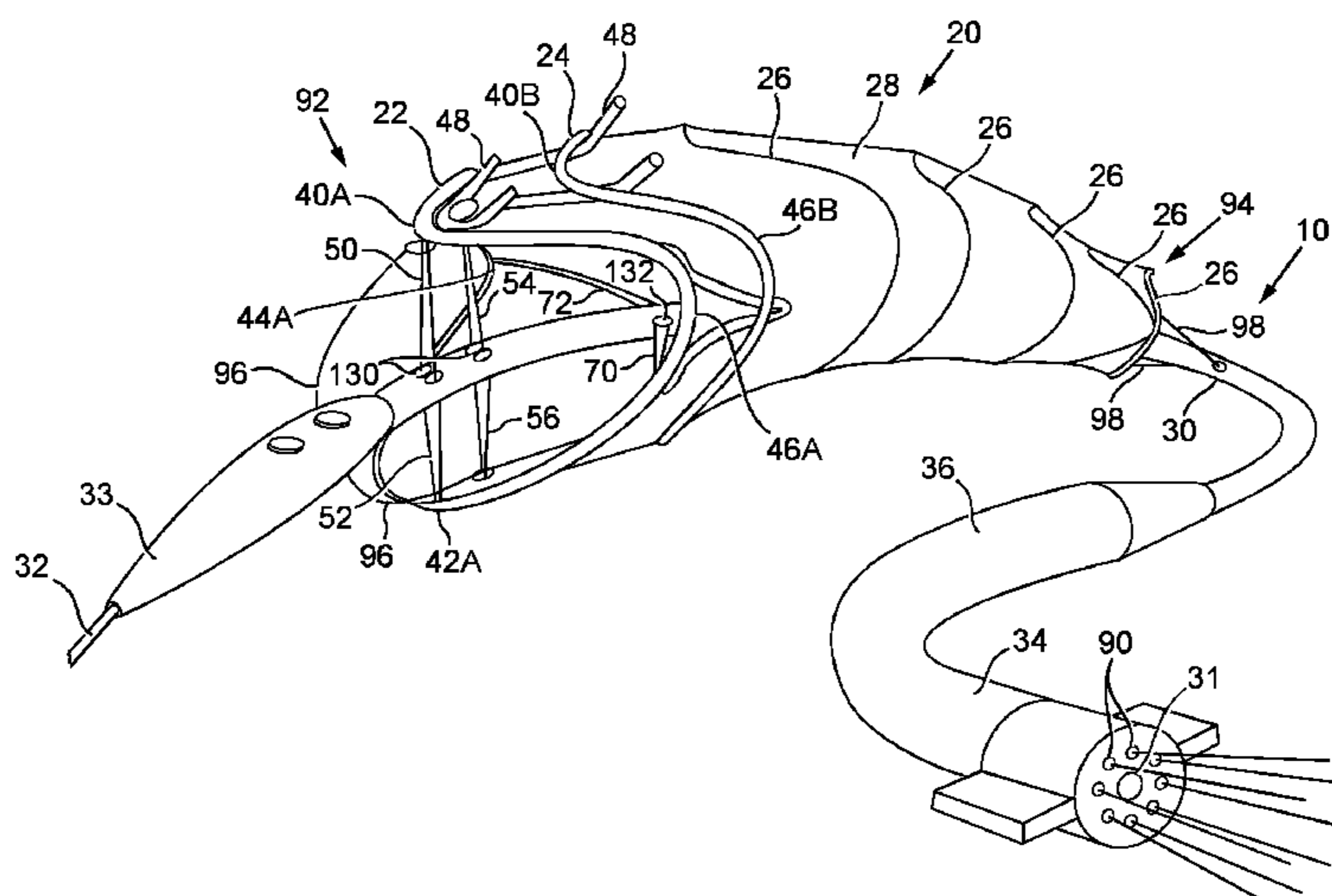
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(57) **ABSTRACT**

A control system for deploying a stent graft (20) including a resilient ring stent (22) at one end, provides a delivery shaft (30) carrying the stent graft with the ring stent adjacent a proximal end (33) of the shaft for insertion into a lumen; a retractable sleeve (36) for containing the stent graft on the delivery shaft; a control handle (38) adjacent the distal end (34), of the delivery shaft; and stent peak (50, 52) and valley (70, 72) control wires extending from the proximal end of the delivery shaft to peak and valley controllers (58; 74, 76) at the control handle. Use of the control wires effects movement and/or rotation of the graft allowing fine control of deployment in the lumen.

58 Claims, 9 Drawing Sheets



(58) **Field of Classification Search**

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 2002/9534; A61F 2002/9665; A61F
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See application file for complete search history.

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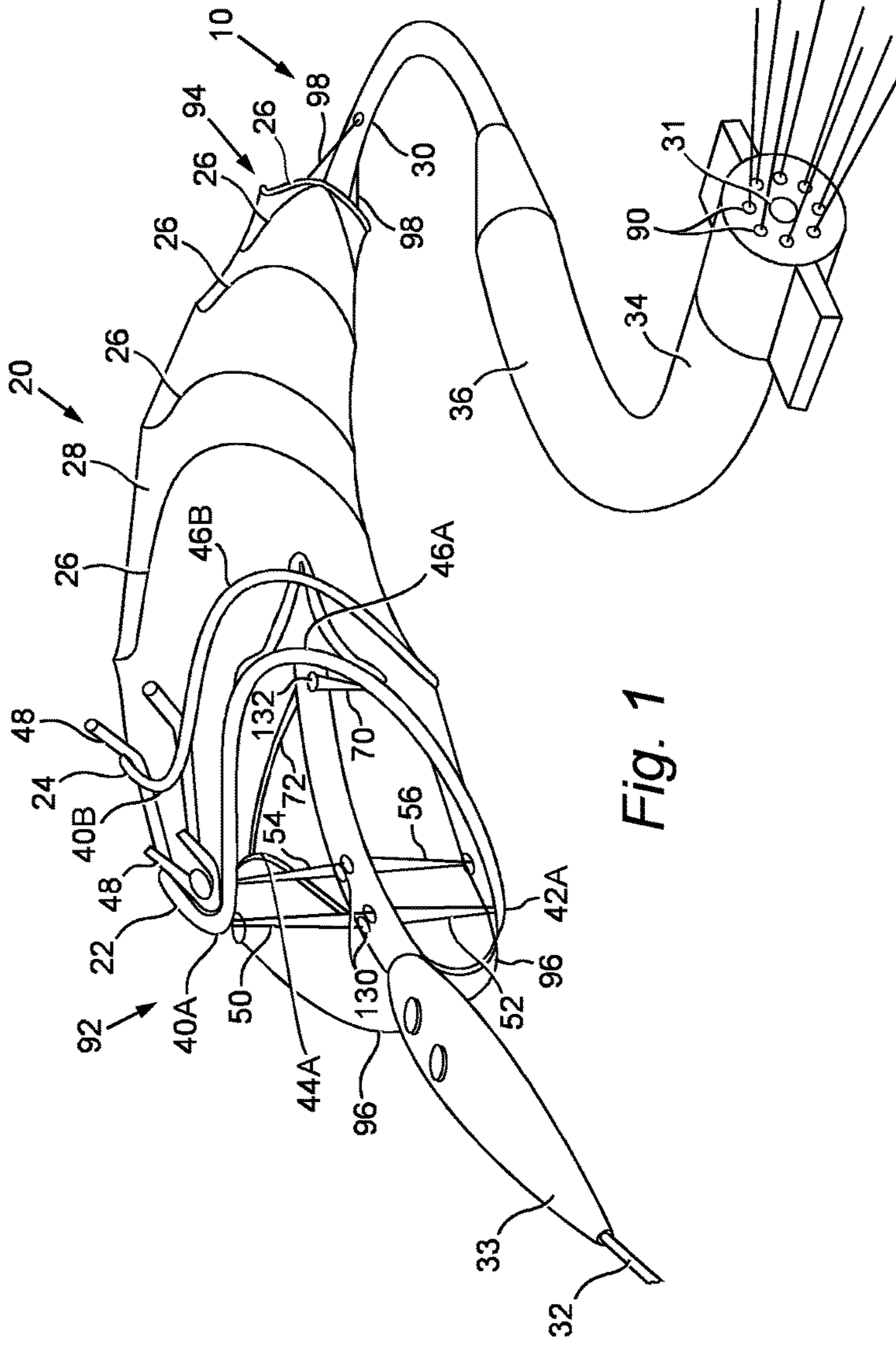


Fig. 1

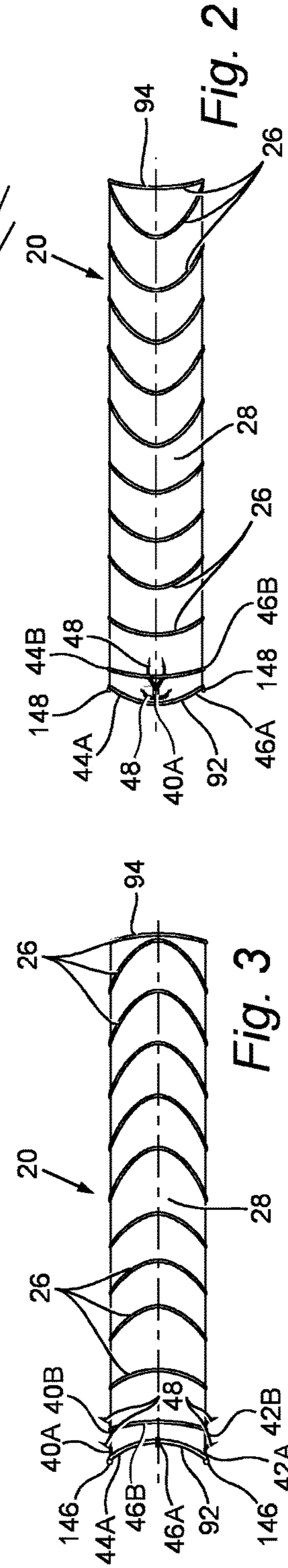


Fig. 2

Fig. 3

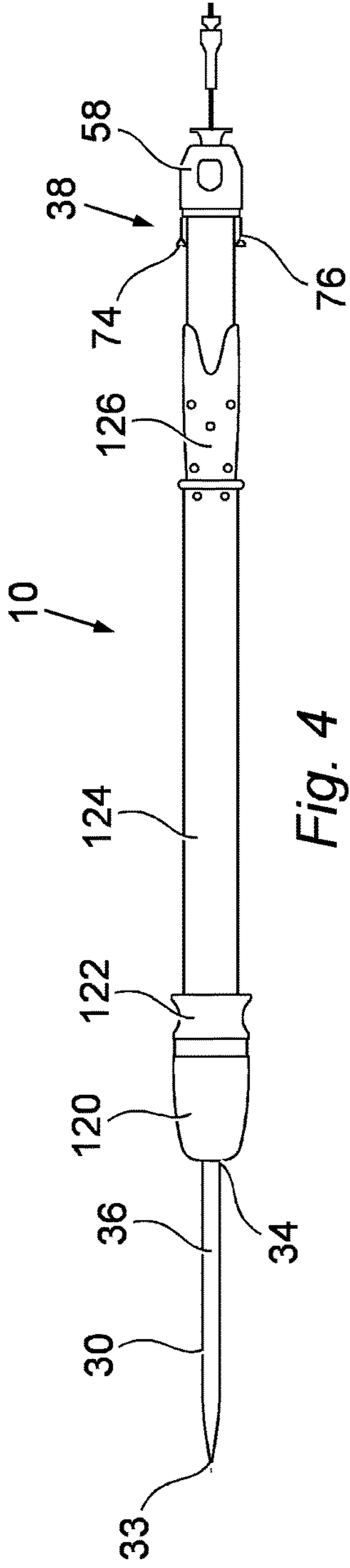


Fig. 4

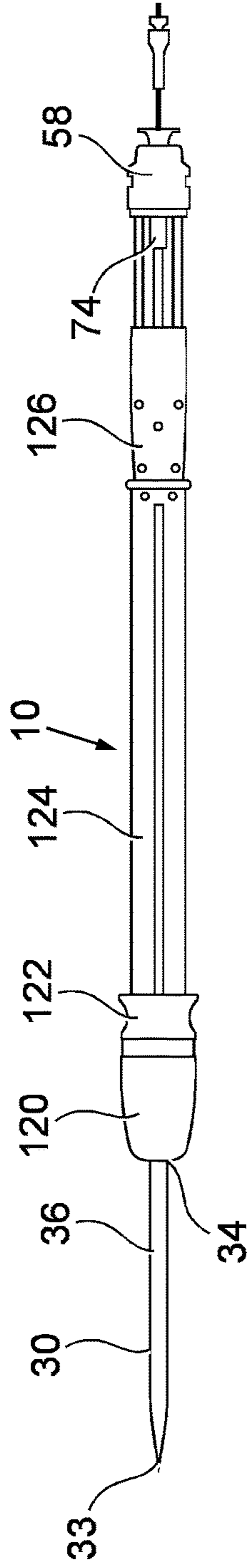


Fig. 5

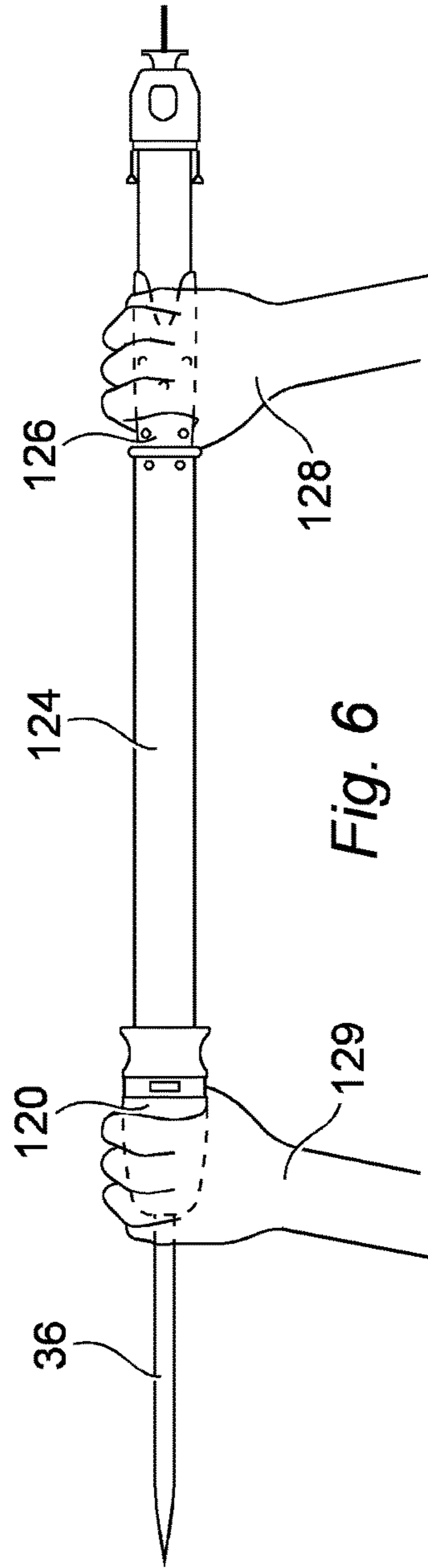


Fig. 6

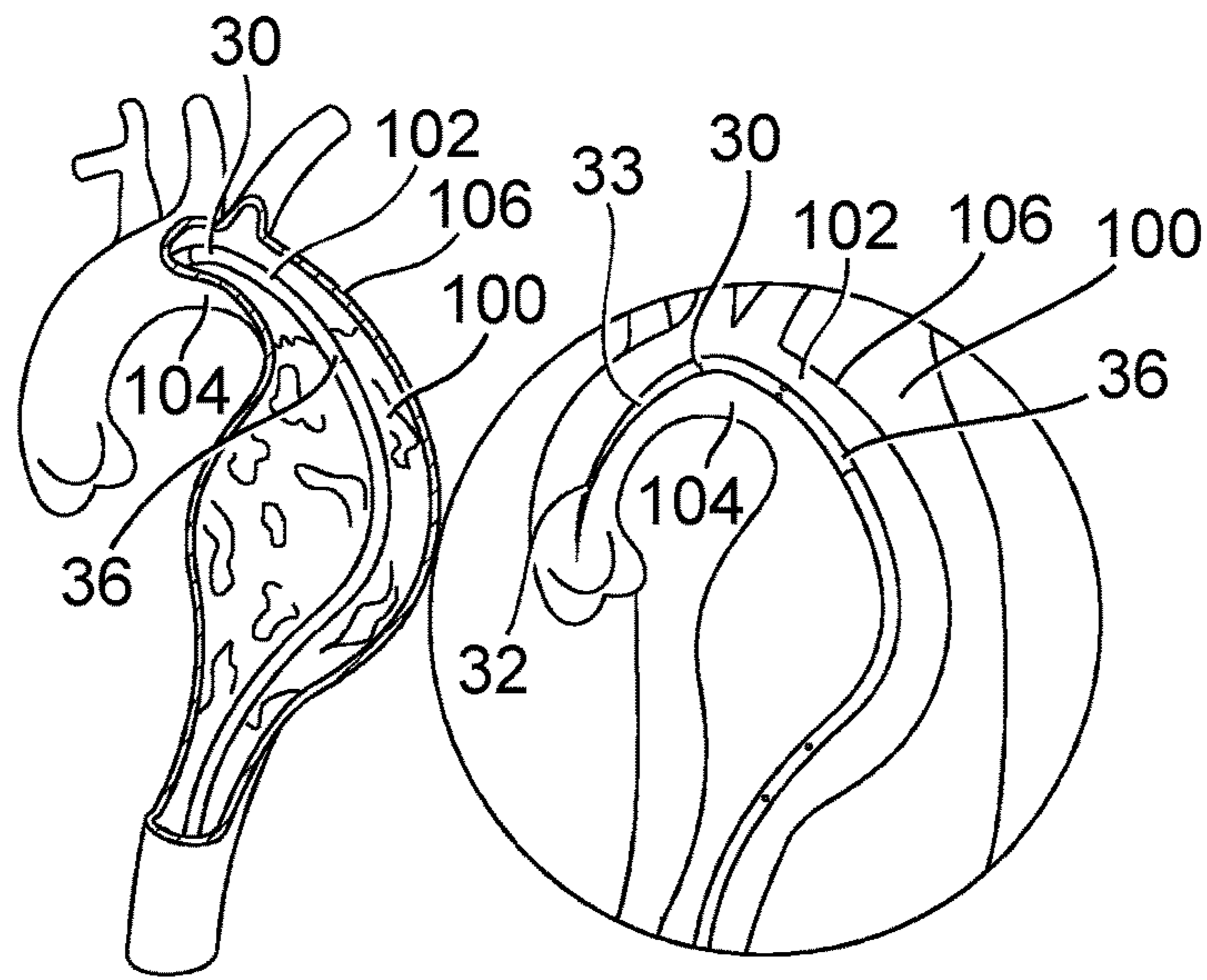


Fig. 7

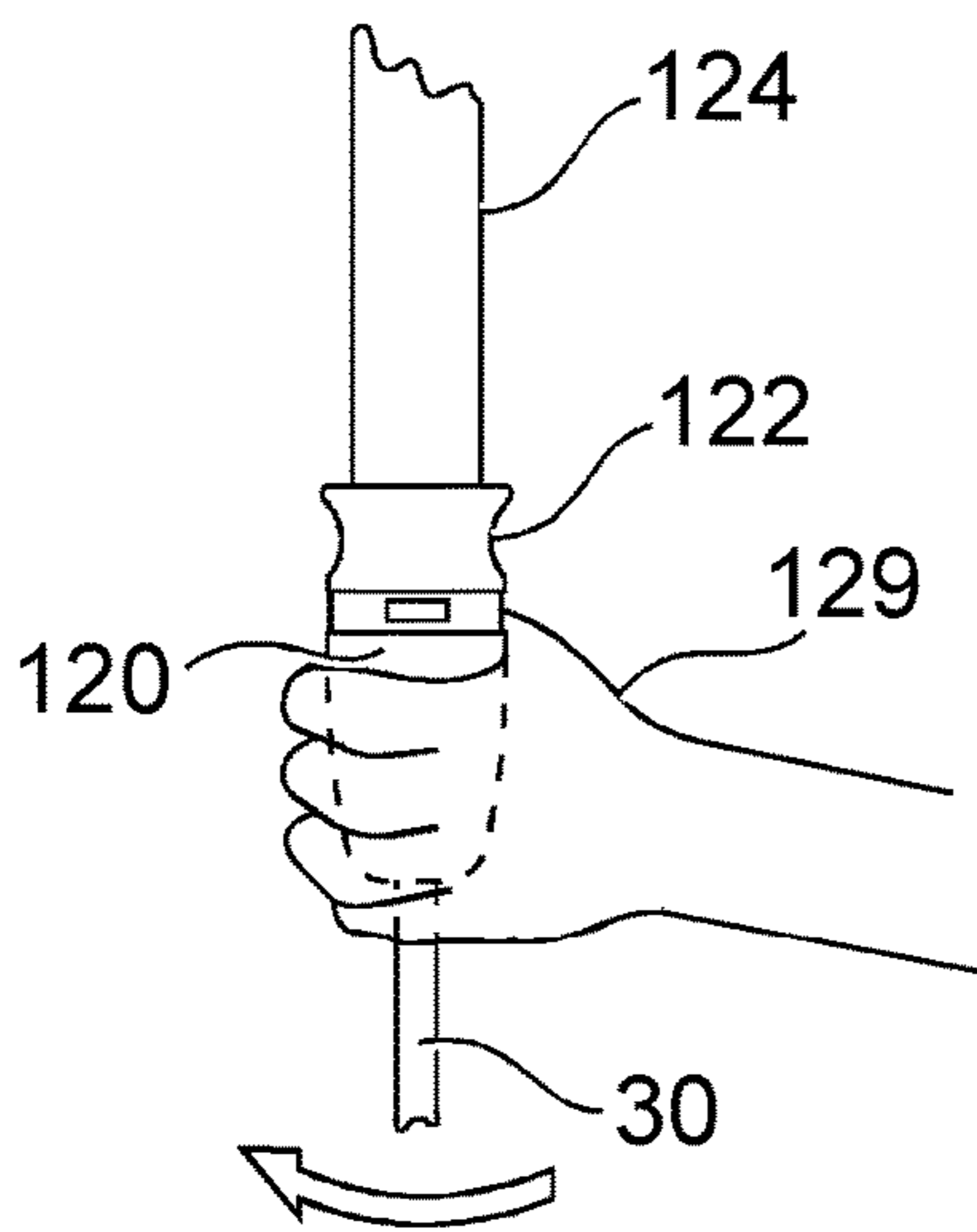


Fig. 8

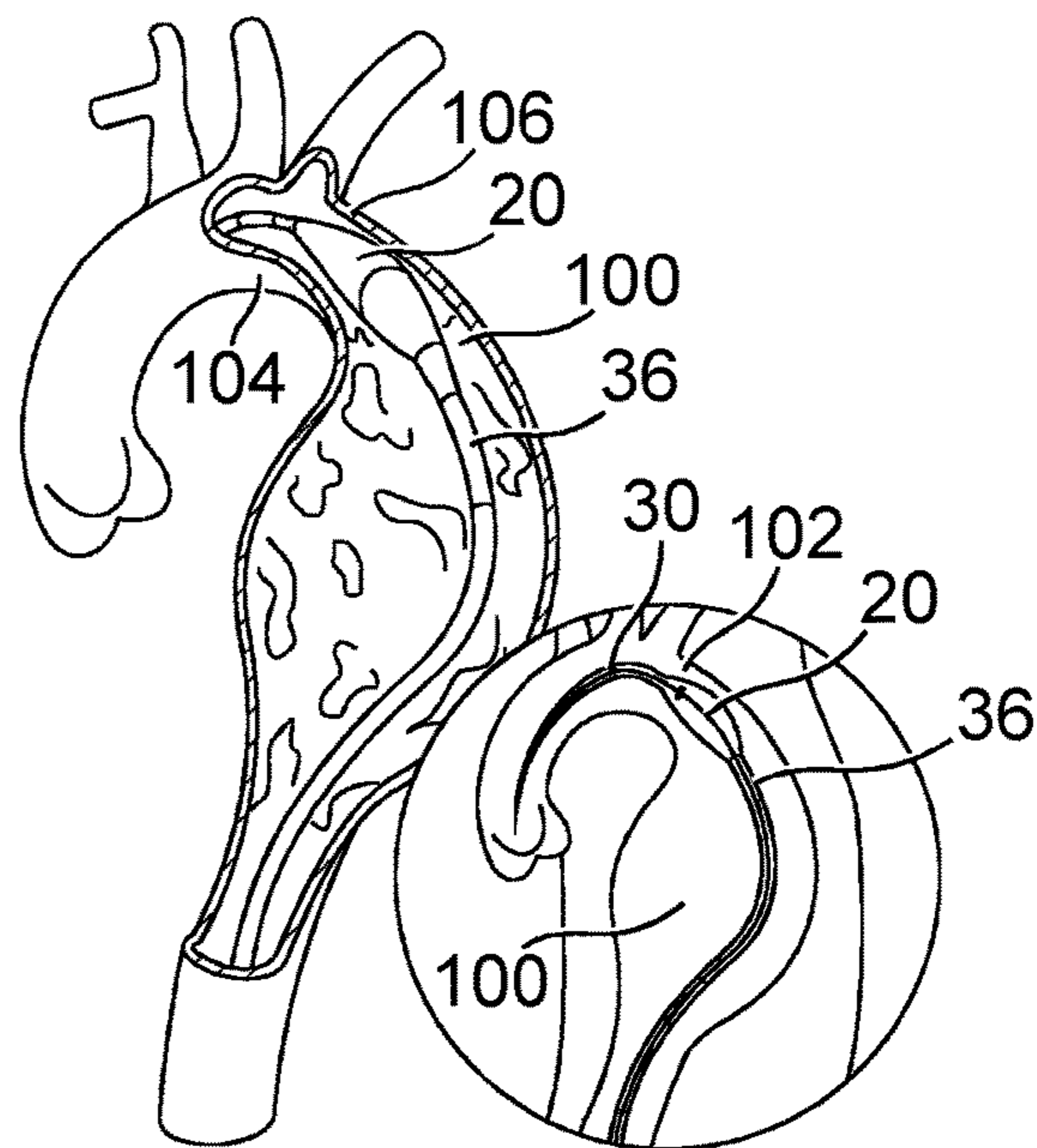


Fig. 9

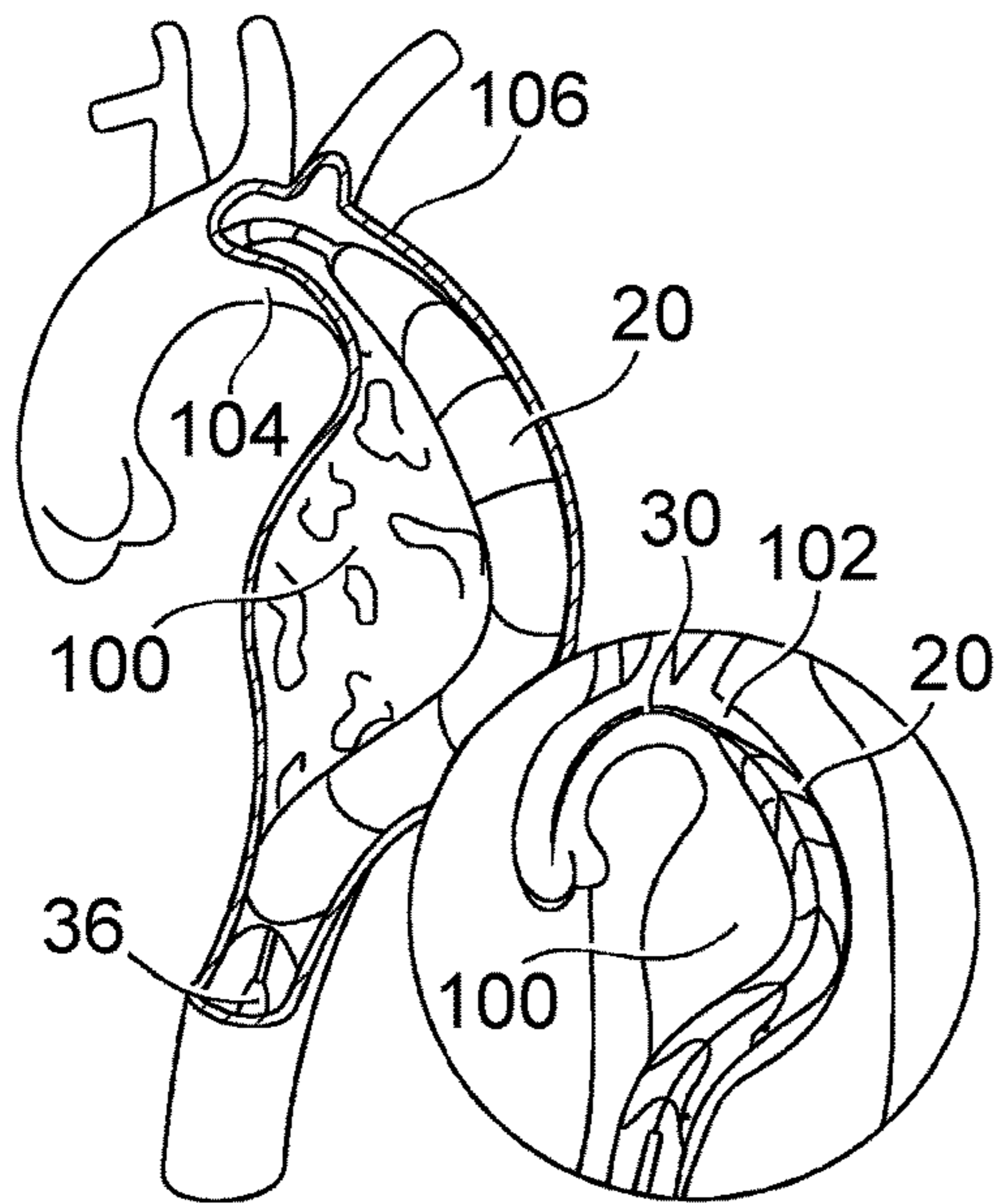


Fig. 10

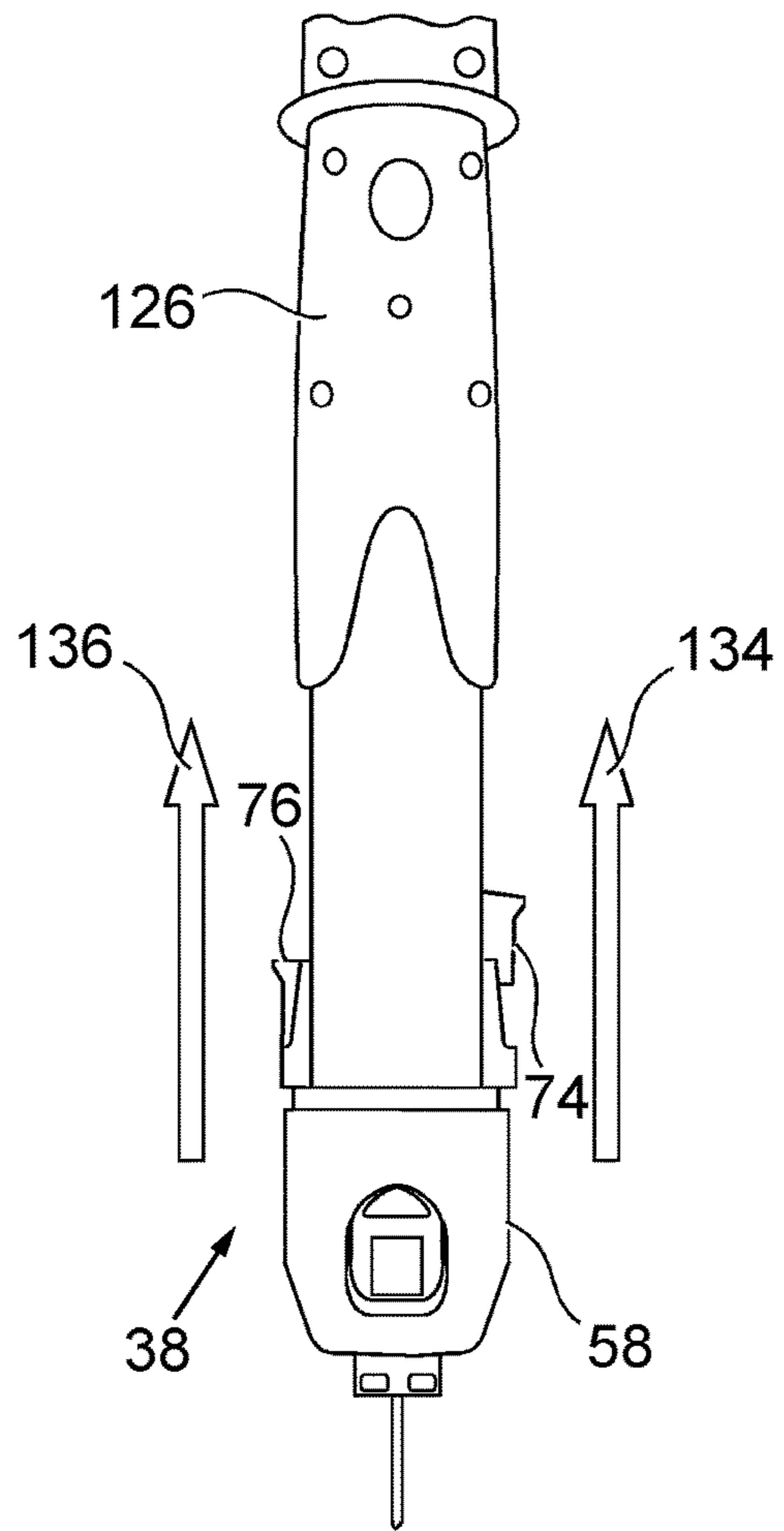


Fig. 11

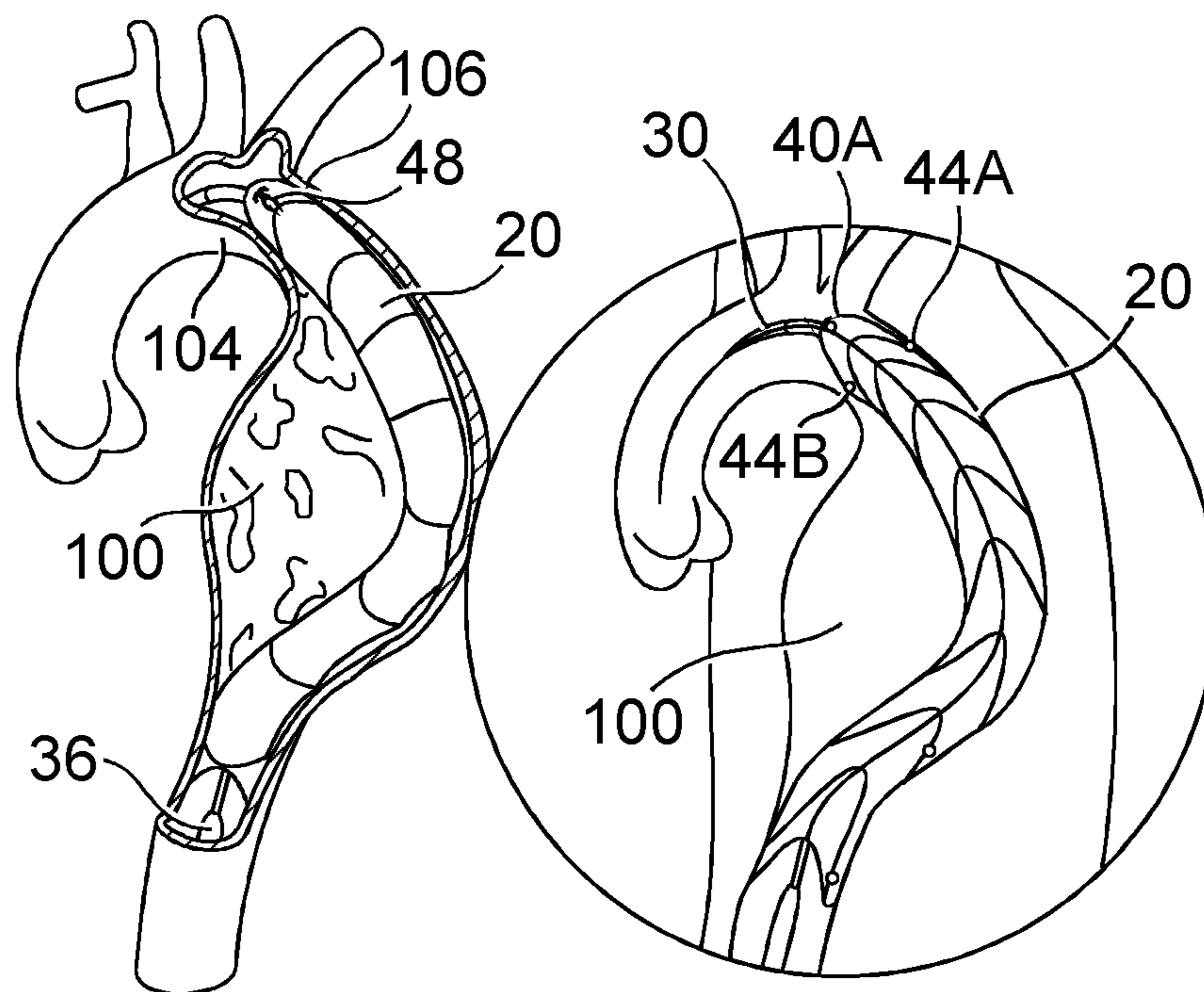


Fig. 13

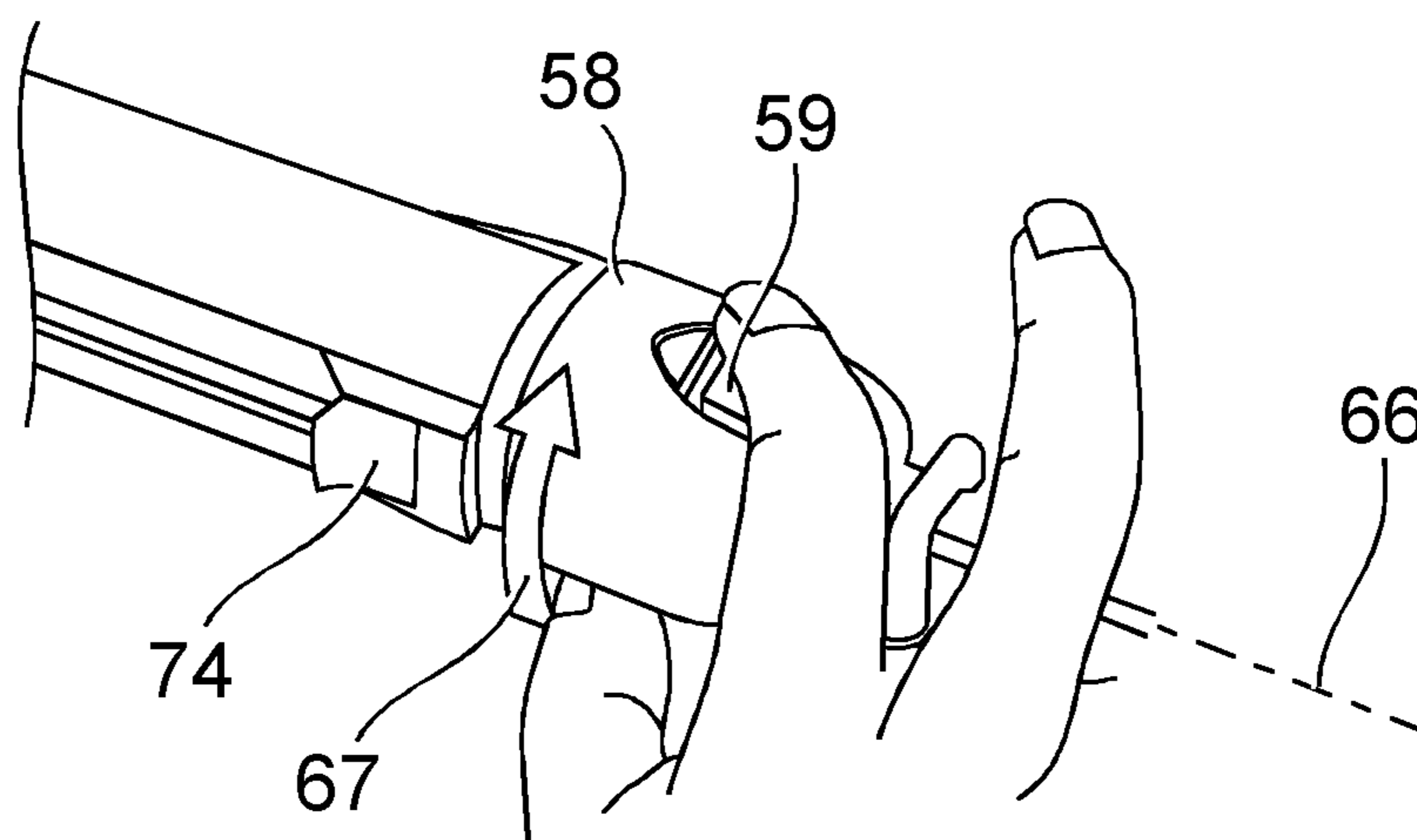


Fig. 12

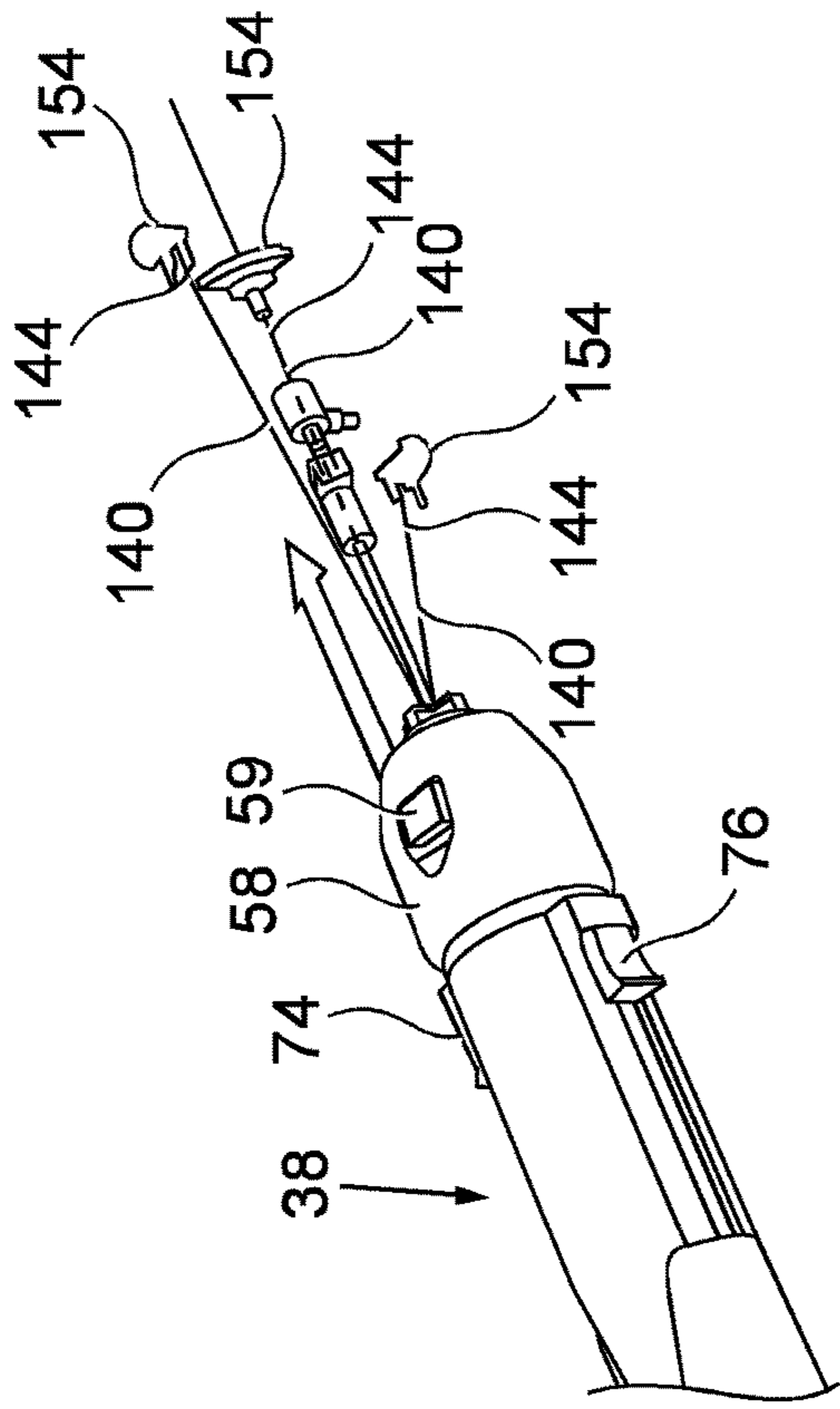


Fig. 14

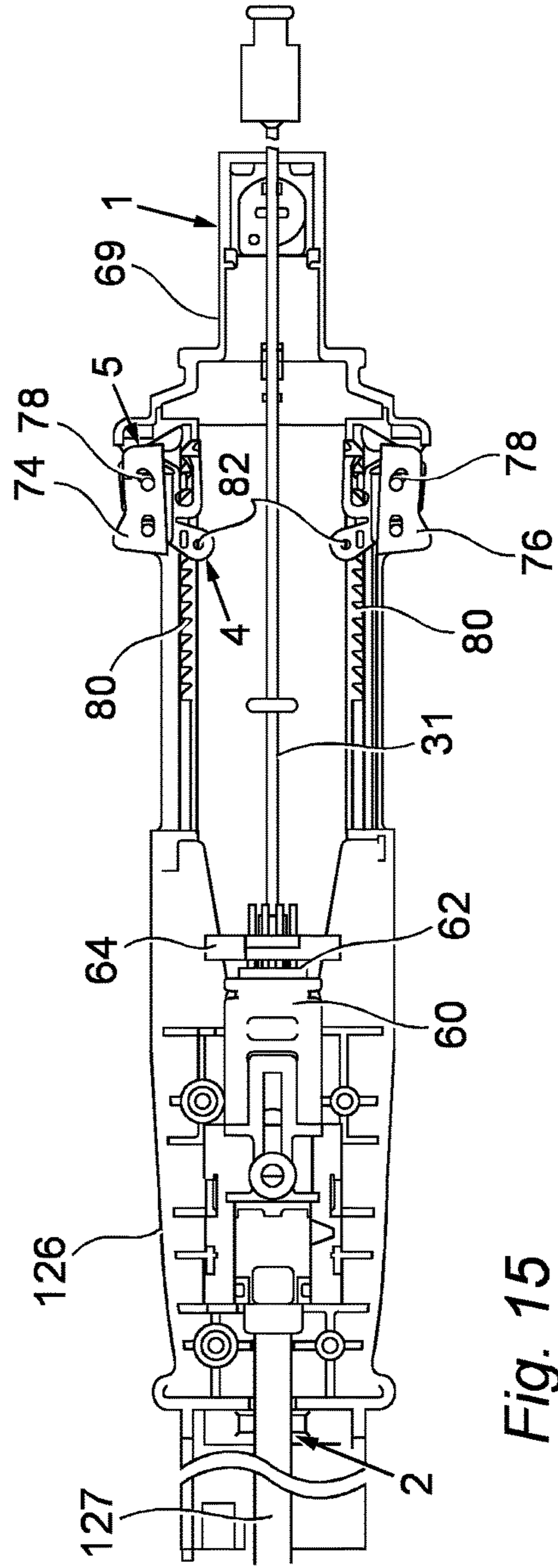
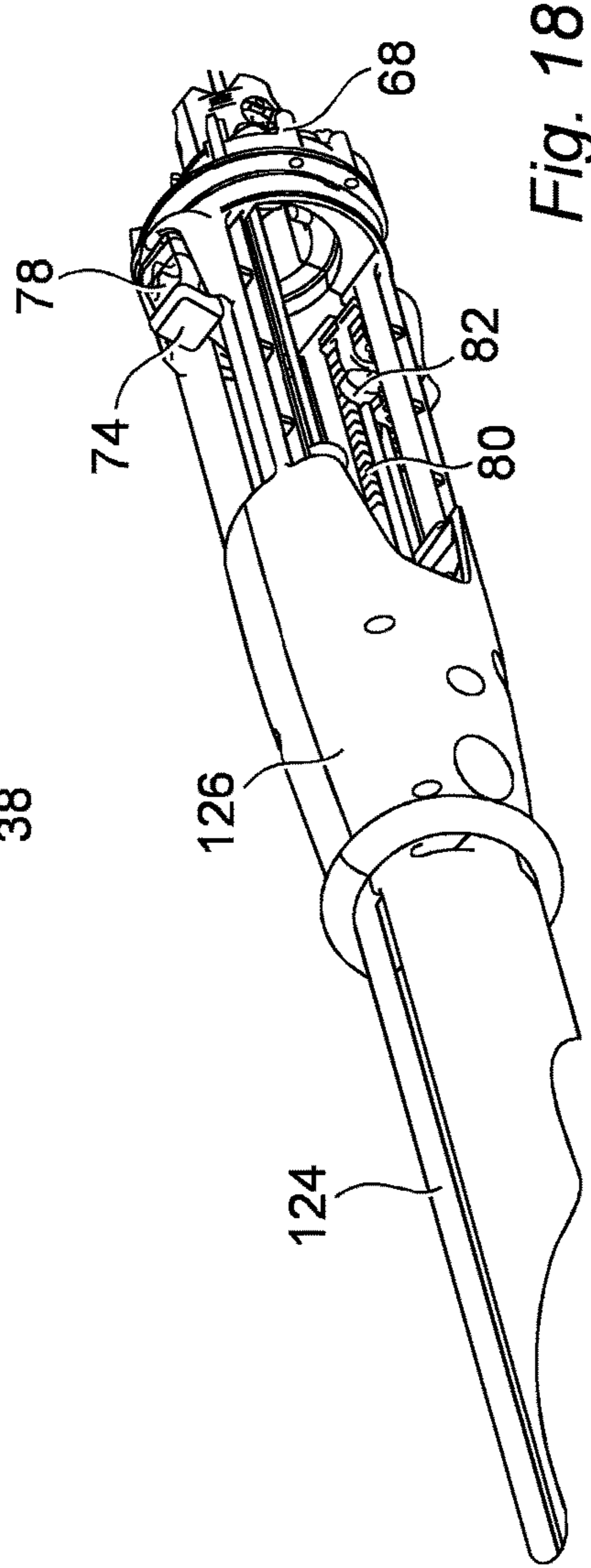
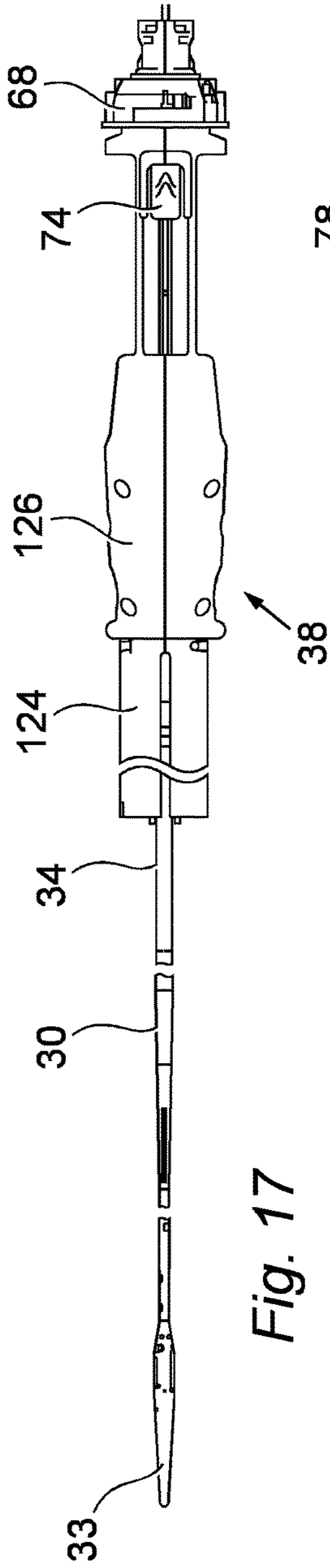
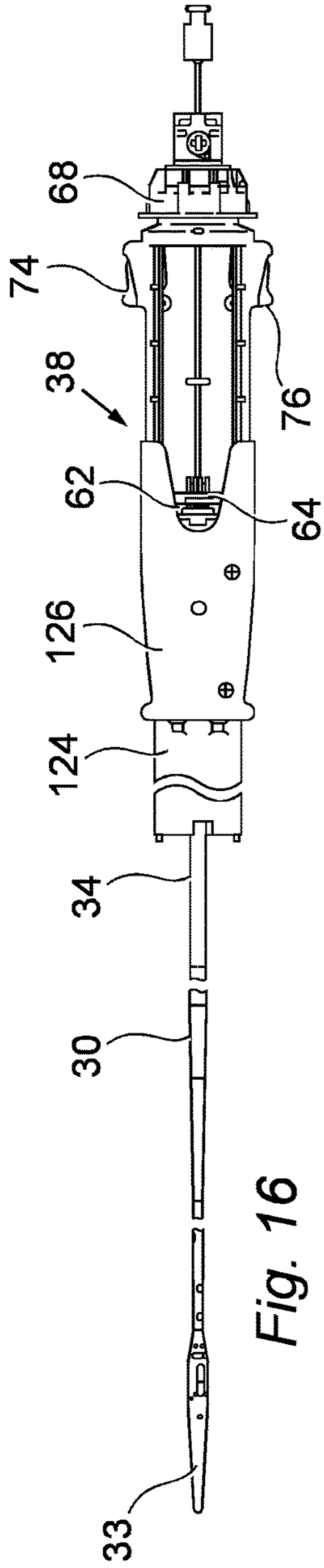


Fig. 15



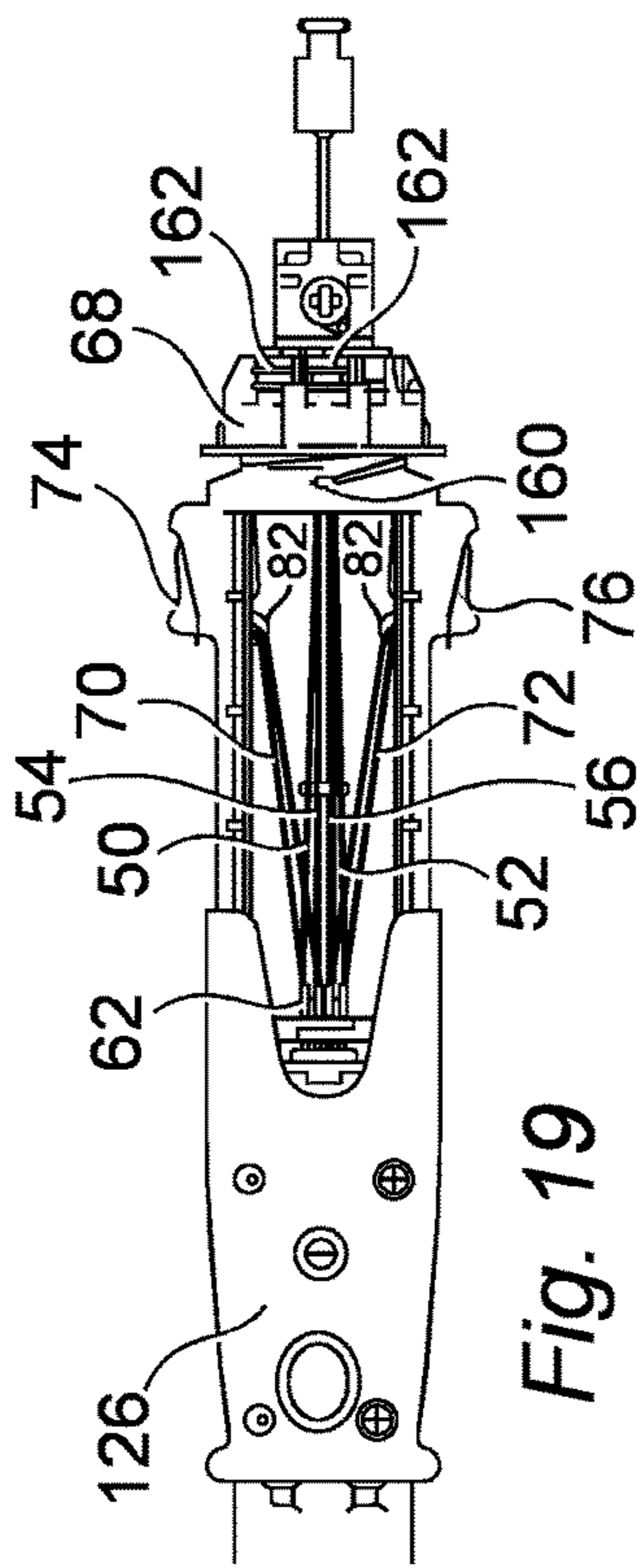


Fig. 19

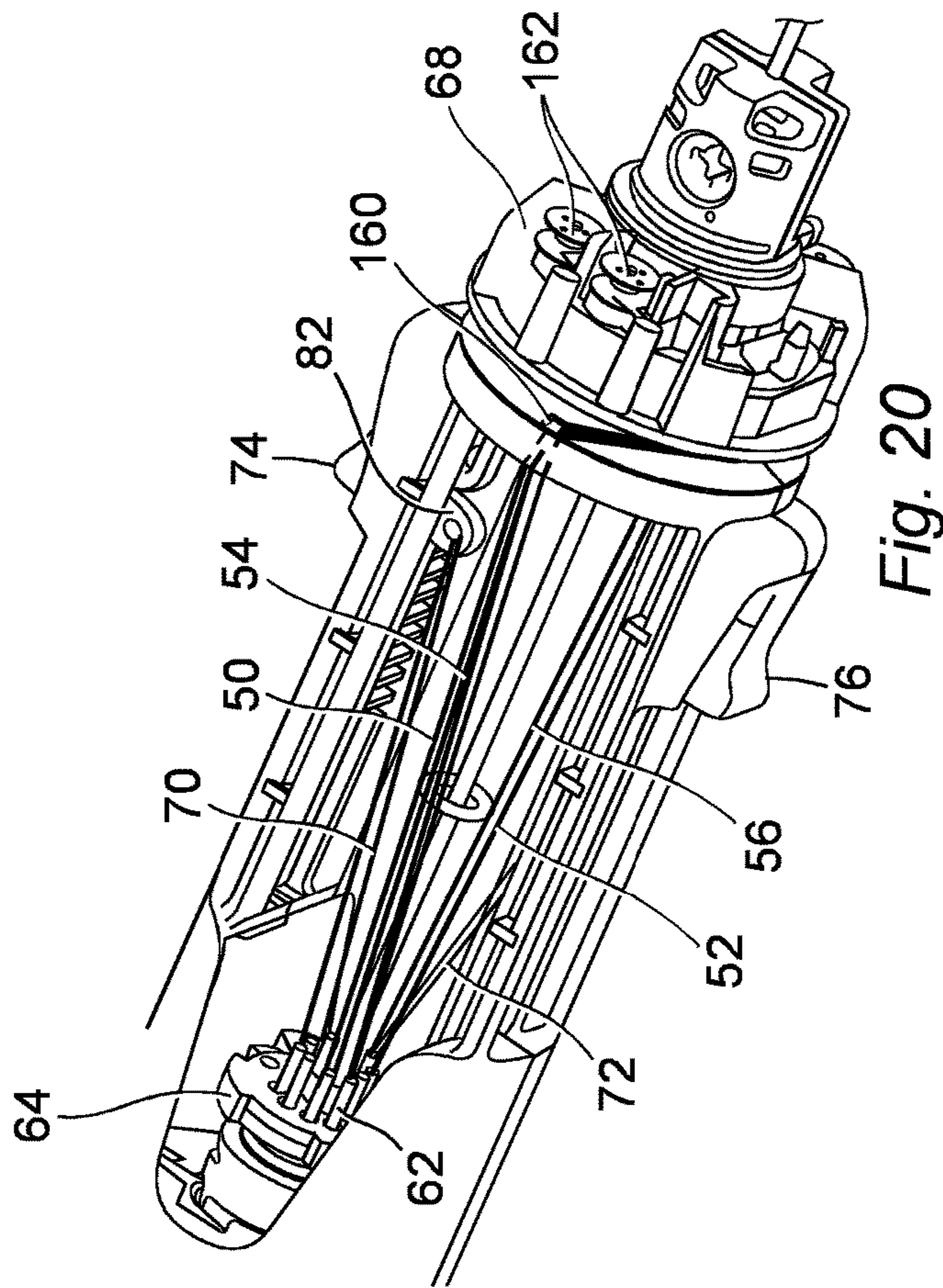


Fig. 20

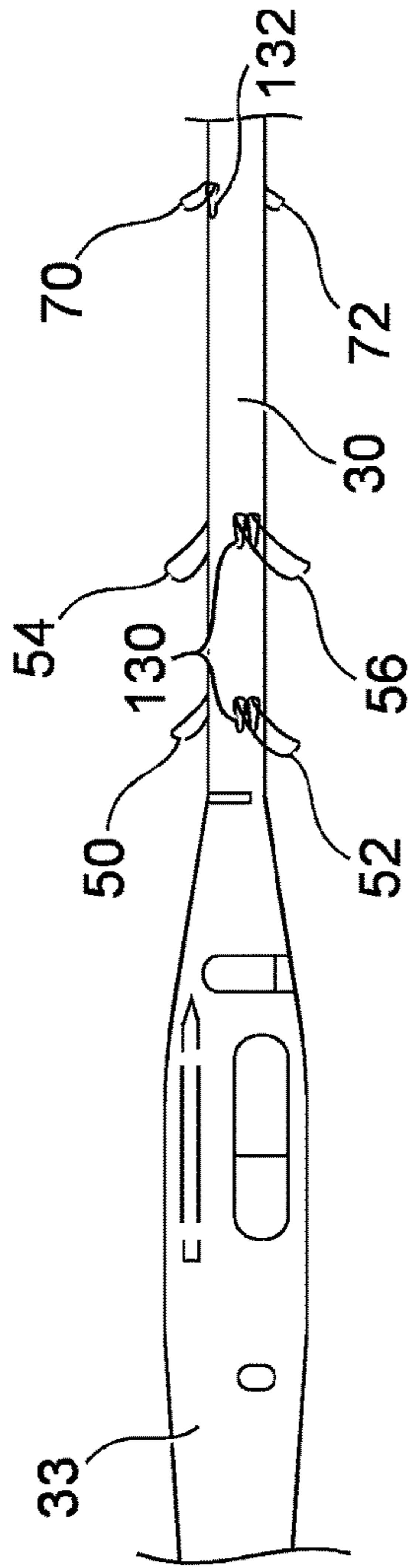


Fig. 21

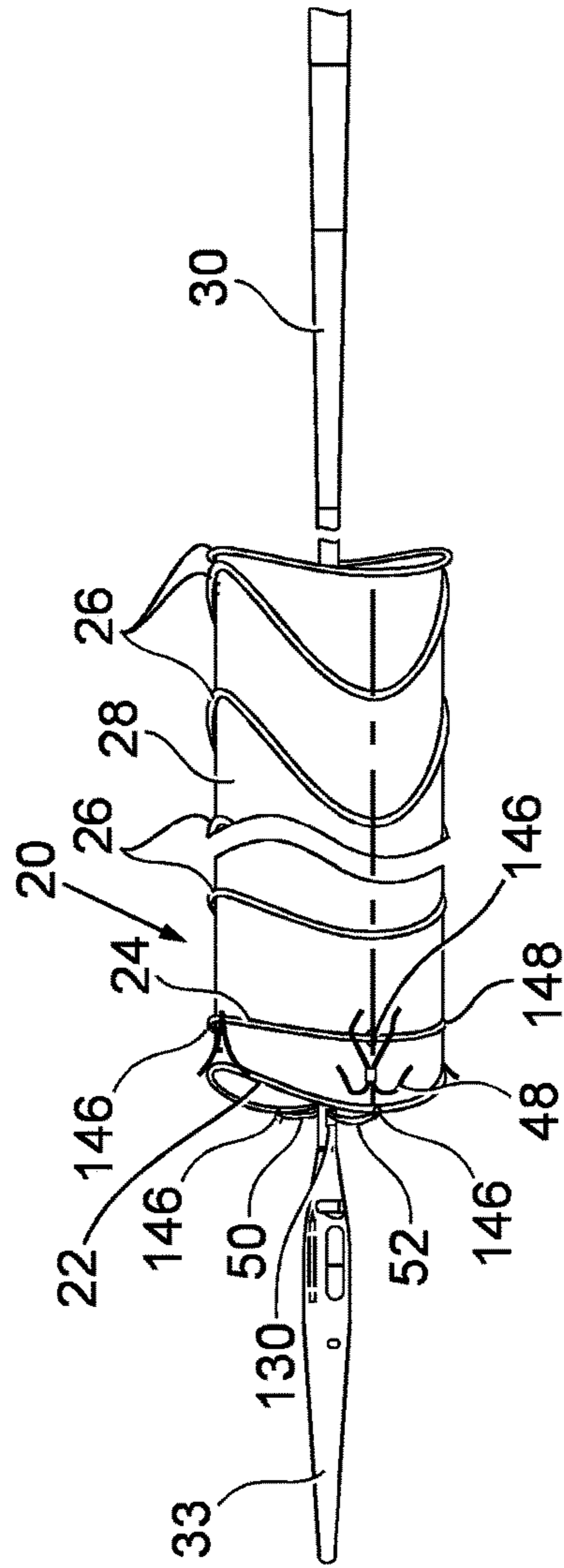


Fig. 22

METHOD AND APPARATUS FOR CONTROLLING THE DEPLOYMENT OF A STENT

This application is a 371 of international application PCT/GB2012/051235, filed Jun. 1, 2012, and claims priority from GB application 1109305.1, filed Jun. 3, 2011, which is incorporated herein by reference in its entirety.

FIELD OF THE INVENTION

The present invention relates to the deployment of a stent element, and in particular to the deployment of a stent element to stabilise a tubular prosthesis.

BACKGROUND TO THE INVENTION

Artificial prostheses consisting of a tubular conduit having an open lumen are well-known and are used in medicine to replace diseased or damaged natural body lumens, such as, for example, blood vessels or other hollow organs for example bile ducts, sections of intestine or the like. The most common use of such artificial prostheses is to replace diseased or damaged blood vessels.

A number of vascular disorders can be treated by use of an artificial prosthesis. One relatively common vascular disorder is an aneurysm. Aneurysm occurs when a section of natural blood vessel wall, typically of the aortic artery, dilates and balloons outwardly. Whilst small aneurysms cause little or no symptoms, larger aneurysms pose significant danger to a patient. Rupture of an aortic aneurysm can occur without warning and is usually fatal, so significant emphasis is placed on early diagnosis and treatment. With an increasing ageing population, the incidence of aneurysm continues to rise in western societies.

Provided that an aneurysm is diagnosed prior to rupture, surgical treatment to repair the affected vessel wall is effective. Surgical treatment of aneurysm involves the replacement or reinforcement of the aneurysmal section of aorta with a synthetic graft or prosthesis under general anaesthesia allowing the patient's abdomen or thorax to be opened (see Parodi et al., *Annals of Vascular Surgery* (1991) 5:491-499). The patient will then have a normal life expectancy.

Surgical repair of aneurysm is however a major and invasive undertaking and there has been much effort in developing less invasive methods. Currently, aneurysm repair generally involves the delivery by catheter of a fabric or ePTFE graft which is retained at the required location by deployment of metallic stent elements. The ability to deliver the graft/stent device by catheter reduces the surgical intervention to a small cut-down to expose the femoral artery and, in suitable circumstances, the device can be deployed percutaneously. Catheter delivery is beneficial since the reduced invasive nature of the procedure allows utilisation of a local anaesthetic and leads to reduced mortality and morbidity, as well as decreased recovery time. For example, endovascular repair is typically used for repair of infra-renal abdominal aortic aneurysms where the graft is placed below the renal arteries. Many different types of devices useful for endovascular repair are now available, for example a resiliently engaging endovascular element described in U.S. Pat. No. 6,635,080 (Vascutek) or a tubular fabric liner having a radially expandable supporting frame and a radiopaque marker element stitched to the liner as disclosed in U.S. Pat. No. 6,203,568 (Medtronic).

However, whilst the endovascular repair of aneurysms is now accepted as the method of choice, the technique has significant limitations and is not suitable for all patients.

As mentioned above, other vascular disorders are treatable by use of a vascular prosthesis. Examples include (but not limited to) occlusions, stenosis, vascular damage due to accident or trauma, and the like. Vascular prostheses are also used in by-pass techniques.

Endovascular techniques involve the delivery of a prosthesis by catheter. Since the internal lumen of the catheter defines the maximum dimensions of the prosthesis to be inserted, much effort has been expended in the design of prostheses which can be packaged in a minimal volume, and are easy to deploy once positioned at the required location.

One successful type of prosthesis is a stent graft comprising a conduit formed of a flexible sleeve attached to a rigid support or stent. The sleeve will typically be made of a fabric (usually a knitted or woven fabric) of ePTFE, PTFE, polyester (for example DACRON), polyethylene or polypropylene and may optionally be coated to reduce friction; discourage clotting or to deliver a pharmaceutical agent. The fabric will generally be porous on at least one surface to enable cell ingrowth.

The stent may be self-expandable and formed of a shape memory material, such as nitinol (a nickel-titanium alloy).

The stent grafts are inserted using a delivery catheter and, once correctly located at the site requiring treatment, are deployed by the withdrawal of a delivery sheath of the delivery catheter. The self-expandable stents are deployed by expanding radially upon release from the delivery sheath. Once deployed, the stents hold the graft in location by contact with the inner wall of the blood vessel.

One suitable stent design is a series of ring stent elements formed from discrete rings of a shape memory material, such as nitinol, attached to the fabric sleeve at spaced intervals. Such a design fulfils the requirements for minimal volume when packaged and, once delivered, readily expands to maintain the patency of the fabric lumen. However, stent grafts having such ring stent elements have the disadvantage that the rings are not readily capable of being contracted again and so do not allow adjustment of the position of the stent graft once deployed.

It is an object of the present invention to overcome one or more of the aforementioned problems.

SUMMARY OF THE INVENTION

According to a first aspect of the present invention there is provided a delivery apparatus for controlling the deployment of a stent, the apparatus comprising:

- a delivery shaft adapted to carry a tubular prosthesis thereon and having a proximal end for insertion into a lumen and a distal end remote from the proximal end;
- a retractable sleeve adapted to contain a tubular prosthesis carried on the delivery shaft;
- a control handle adjacent the distal end of the delivery shaft;
- at least one stent peak control wire extending from the proximal end of the delivery shaft to the control handle;
- a peak controller at the control handle adapted to control the position of the stent peak control wire.

The term "wire" as used in this specification includes metallic and non-metallic wire, thread and filament.

The apparatus may comprise two or more stent peak control wires extending from the proximal end of the

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delivery shaft to the control handle, and the peak controller may be adapted to control the position of the two or more stent peak control wires.

The apparatus may further comprise:

at least one stent valley control wire extending from the proximal end of the delivery shaft to the control handle; and

a valley controller at the control handle adapted to control the position of the stent valley control wire.

The apparatus may comprise two stent valley control wires extending from the proximal end of the delivery shaft to the control handle, and the apparatus may include two valley controllers at the control handle, each valley controller being adapted to control the position of a respective stent valley control wire.

The delivery shaft may include a plurality of longitudinal passages, each control wire extending through a respective longitudinal passage.

The apparatus may include one or more tow wires provided at the proximal end of the delivery shaft adapted for connection to a proximal end of a tubular prosthesis carried on the delivery shaft. The apparatus may include one or more back tether wires provided between the proximal end of the delivery shaft and the control handle adapted for connection to a distal end of a tubular prosthesis carried on the delivery shaft.

The control handle may include a manifold having a plurality of apertures, each control wire extending through a respective aperture. The manifold may include a plurality of haemostasis valves switchable between open and closed positions.

The peak controller may comprise a rotatable collar. The rotatable collar may be mounted for rotation about a longitudinal axis of the control handle. The peak controller may include a winder which rotates with the collar and on which are wound the stent peak control wires.

Each valley controller may comprise a slider. Each valley controller may include an actuating surface and a ratchet adapted to allow the slider to move when the actuating surface is depressed and to hold the slider in an engaged position when the actuating surface is released. The sliders may be mounted for longitudinal sliding movement. Each slider may include a wire connector to which is connected the respective stent valley control wire.

Preferably the delivery shaft includes a plurality of peak apertures adjacent the proximal end thereof, each stent peak control wire extending through one of said plurality of peak apertures. Preferably the delivery shaft includes a plurality of valley apertures adjacent the proximal end thereof, each stent valley control wire extending through one of said plurality of valley apertures. Preferably the valley apertures are arranged between the peak apertures and the distal end.

The control wires may be looped wires which extend from the control handle to the proximal end of the delivery shaft and back to the control handle.

The apparatus may include a plurality of release wires extending from the control handle along the delivery shaft and having a proximal end at or near the proximal end of the delivery shaft and a distal end at the control handle. The proximal end of each release wire is adapted to be connected to a respective connection between a tubular prosthesis carried on the delivery shaft and one of the stent peak control wires, the stent valley control wires, the tow wires and the back tether wires. The distal end of each release wire may be connected to a release clip mounted on the control handle. The release clip may be removed from the control handle and pulled with its corresponding release wire or wires away

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from the apparatus, thereby releasing the respective connection between a tubular prosthesis carried on the delivery shaft and one of the stent peak control wires, the stent valley control wires, the tow wires and the back tether wires.

According to a second aspect of the present invention there is provided a method of deploying a ring stent, the method comprising the steps of:

providing a tubular prosthesis including a ring stent thereon on a delivery shaft, the delivery shaft having a proximal end for insertion into a lumen and a distal end remote from the proximal end, and having two stent peak control wires extending from the proximal end to the distal end;

attaching the first ends of two stent peak control wires to first and second diametrically opposed peak connection points on the tubular prosthesis adjacent the ring stent; attaching the second ends of the two stent peak control wires to a peak controller provided at a control handle adjacent the distal end of the delivery shaft;

holding the two stent peak control wires under tension such that the peak connection points are held by the stent peak control wires in a folded arrangement towards the delivery shaft; and

operating the peak controller to permit movement of the stent peak control wires along the delivery shaft such that the peak connection points are urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent adopts a partially deployed position.

The peak connection points may be on the ring stent.

The delivery shaft may include two stent valley control wires extending from the proximal end to the distal end, and the method may comprise the further steps of:

attaching the first ends of the two stent valley control wires to first and second diametrically opposed valley connection points on the tubular prosthesis adjacent the ring stent, the valley connection points being provided midway between the peak connection points;

attaching the second ends of the two stent valley control wires to first and second valley controllers respectively provided at the control handle;

holding the two valley peak control wires under tension such that the valley connection points are held by the stent valley control wires in a folded arrangement towards the delivery shaft; and

selectively operating one of the first and second valley controllers to permit movement of the stent valley control wires along the delivery shaft such that the first or second valley connection point is urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent is rotated in the partially deployed position.

The valley connection points may be on the ring stent or may be on a valley line between two adjacent ring stents on the tubular prosthesis.

The method may include the further step of selectively operating the other of the first and second valley controllers to permit movement of the stent valley control wires along the delivery shaft such that both the first and second valley connection points are urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent adopts a fully deployed position.

The method may include the further step of selectively operating one or more of the peak controller and first and second valley controllers to pull at least some of the control wires along the delivery shaft such that one or more of the connection points adjacent the ring stent are moved towards

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the delivery shaft against the resilience of the ring stent to permit the ring stent to be moved in position.

The control wires may be looped wires which extend from the control handle to the peak or valley connection points and back to the control handle. The peak or valley connection points may be eyelets through which the looped wires are threaded.

The step of operating the peak controller may include rotating a rotatable collar to wind or unwind the stent peak control wires on a winder.

The step of operating the first or second valley controller may include moving a slider which includes a wire connector to which is connected the respective stent valley control wire.

According to a third aspect of the present invention there is provided a method of causing the rotation of a ring stent in a lumen, the method comprising the steps of:

providing a tubular prosthesis including a ring stent thereon on a delivery shaft, the delivery shaft having a proximal end for insertion into a lumen and a distal end remote from the proximal end, and having two stent valley control wires extending from the proximal end to the distal end;

attaching the first ends of two stent valley control wires to first and second diametrically opposed valley connection points on the tubular prosthesis adjacent the ring stent;

attaching the second ends of the two stent valley control wires to first and second valley controllers respectively provided at a control handle adjacent the distal end of the delivery shaft;

holding the two stent valley control wires under tension such that the valley connection points are held by the stent valley control wires in a folded arrangement towards the delivery shaft; and

selectively operating one of the first and second valley controllers to permit movement of the stent valley control wires along the delivery shaft such that the first or second valley connection point is urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent is rotated.

The valley connection points may be on the ring stent or may be on a valley line between two adjacent ring stents on the tubular prosthesis.

The method may include the further step of selectively operating the other of the first and second valley controllers to permit movement of the stent valley control wires along the delivery shaft such that both the first and second valley connection points are urged by the resilience of the ring stent to move away from the delivery shaft by a respective selected amount and the ring stent adopts a selected rotational position.

The delivery shaft may include two stent peak control wires extending from the proximal end to the distal end, and the method may comprise the further steps of:

attaching the first ends of the two stent peak control wires to first and second diametrically opposed peak connection points on the tubular prosthesis adjacent the ring stent, the peak connection points being provided midway between the valley connection points;

attaching the second ends of the two stent peak control wires to a peak controller provided at a control handle adjacent the distal end of the delivery shaft;

holding the two stent peak control wires under tension such that the peak connection points are held by the stent peak control wires in a folded arrangement towards the delivery shaft; and

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operating the peak controller to permit movement of the stent peak control wires along the delivery shaft such that the peak connection points are urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent adopts a fully deployed position.

The peak connection points may be on the ring stent.

The method may include the further step of selectively operating one or more of the peak controller and first and second valley controllers to pull at least some of the control wires along the delivery shaft such that one or more of the connection points adjacent the ring stent are moved towards the delivery shaft against the resilience of the ring stent to permit the ring stent to be moved in position.

The control wires may be looped wires which extend from the control handle to the peak or valley connection points and back to the control handle. The peak or valley connection points may be eyelets through which the looped wires are threaded.

The step of operating the peak controller may include rotating a rotatable collar to wind or unwind the stent peak control wires on a winder.

The step of operating the first or second valley controller may include moving a slider which includes a wire connector to which is connected the respective stent valley control wire.

According to a fourth aspect of the present invention there is provided a stent system comprising a tubular prosthesis including a resilient ring stent at a first end thereof, and an apparatus for controlling the deployment of the ring stent, the apparatus comprising:

a delivery shaft carrying the tubular prosthesis thereon and having a proximal end for insertion into a lumen and a distal end remote from the proximal end, the ring stent being adjacent the proximal end;

a retractable sleeve adapted to contain the tubular prosthesis carried on the delivery shaft;

a control handle adjacent the distal end of the delivery shaft;

a first stent peak control wire extending from the proximal end of the delivery shaft to the control handle; and

a peak controller at the control handle adapted to control the position of the first stent peak control wire; wherein the stent peak control wire is connected to a first peak connection point on the tubular prosthesis adjacent the ring stent.

The apparatus may further include a second stent peak control wire extending from the proximal end of the delivery shaft to the control handle, wherein the peak controller is adapted to control the position of the second stent peak control wire, and wherein the second stent peak control wire is connected to a second peak connection point on the tubular prosthesis adjacent the ring stent diametrically opposed to the first peak connection point.

The peak connection points may be on the ring stent.

The apparatus may further include a first stent valley control wire extending from the proximal end of the delivery shaft to the control handle, and a first valley controller adapted to control the position of the first stent valley control wire, wherein the first stent valley control wire is connected to a first valley connection point on the tubular prosthesis adjacent the ring stent interposed between the first and second peak connection points.

The apparatus may further include a second stent valley control wire extending from the proximal end of the delivery shaft to the control handle, and a second valley controller adapted to control the position of the second stent valley control wire, wherein the second stent valley control wire is

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connected to a second valley connection point on the tubular prosthesis adjacent the ring stent diametrically opposed to the first valley connection point.

The valley connection points may be on the ring stent or may be on a valley line between two adjacent ring stents on the tubular prosthesis.

Preferably the apparatus is an apparatus according to the first aspect of the invention.

The ring stent may have the shape of the perimeter of a saddle. The first and second peak connection points may be provided at peaks of the saddle. The first and second valley connection points may be provided at valleys of the saddle. Preferably the distance between the valley connection points and the control handle is less than the distance between the peak connection points and the control handle.

Preferred or alternative features of each aspect or embodiment of the invention apply mutatis mutandis to each other aspect or embodiment of the invention, unless the context demands otherwise.

DESCRIPTION OF THE DRAWINGS

The present invention will now be further described by reference to the following figures, in which:

FIG. 1 is a schematic illustration of a delivery shaft and stent graft of a delivery apparatus of the invention;

FIGS. 2 and 3 are a top view and a side view respectively of the stent graft of FIG. 1;

FIGS. 4 and 5 are a top view and a side view respectively of the apparatus of FIG. 1;

FIG. 6 is a schematic illustration of an operator operating the apparatus of FIG. 1;

FIG. 7 is a schematic illustration of a thoracic arch during a first stage of a method of deployment of a ring stent according to the invention;

FIG. 8 is a side view of the handle of the delivery apparatus during a second stage of a method of deployment of a ring stent according to the invention;

FIG. 9 is a schematic illustration of a thoracic arch during the second stage of a method of deployment of a ring stent according to the invention;

FIG. 10 is a schematic illustration of a thoracic arch during a third stage of a method of deployment of a ring stent according to the invention;

FIG. 11 is a side view of the handle of the delivery apparatus during a fourth stage of a method of deployment of a ring stent according to the invention;

FIG. 12 is a view of the handle of the delivery apparatus during a fifth stage of a method of deployment of a ring stent according to the invention;

FIG. 13 is a schematic illustration of a thoracic arch during the fourth and fifth stages of a method of deployment of a ring stent according to the invention;

FIG. 14 is a view of the handle of the delivery apparatus during a sixth stage of a method of deployment of a ring stent according to the invention;

FIG. 15 is a partial sectional view through the control handle of the delivery apparatus of FIG. 1;

FIGS. 16 and 17 are a partial top view and a partial side view respectively of the delivery apparatus of FIG. 1;

FIG. 18 is a partial isometric view of the control handle of the delivery apparatus of FIG. 1;

FIG. 19 is a partial top view of the control handle of the delivery apparatus of FIG. 1 with the control wires installed;

FIG. 20 is a partial isometric view of the control handle of the delivery apparatus of FIG. 1 with the control wires installed;

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FIG. 21 is a schematic partial view of the proximal end of the delivery shaft of the delivery apparatus of FIG. 1 with the control wires installed; and

FIG. 22 is a schematic view of the stent graft on the delivery shaft of the delivery apparatus of FIG. 1 with the control wires installed.

DESCRIPTION OF EMBODIMENTS OF THE INVENTION

Referring to FIG. 1, there is shown a delivery apparatus 10 used to deliver a tubular prosthesis, for example a stent graft 20, to the thoracic arch. The delivery apparatus 10 includes a delivery shaft 30 having a central passage 31 adapted to travel on a guide wire 32 provided in the thoracic arch, so that the shaft 30 can be extended within the body of a patient to the correct location. Such guide wires are known in the art and are not described further.

The stent graft is shown in more detail in FIGS. 2 and 3. The stent graft includes a first ring stent 22, a second ring stent 24 and a number of other ring stents 26. The ring stents 22, 24, 26 are attached to a sleeve 28 of the stent graft 20. The sleeve 28 is a flexible tubular conduit having the ring stents 22, 24, 26 attached thereto to maintain the patency of the sleeve lumen after deployment. The sleeve 28 is a woven or knitted flexible fabric which is generally impervious to fluid such as blood. Suitable materials include polyester, such as Dacron. The sleeve 28 may be coated to reduce blood clotting, to reduce friction or to deliver a medicament. The stent graft 20 shown in FIGS. 1 to 3 in the expanded or deployed configuration.

The ring stents 22, 24, 26 are typically formed from a resilient alloy material suitable for medical use, such as nitinol, and are generally saddle shaped. The first ring stent 22 has two diametrically opposed ring stent peaks 40A, 42A and two diametrically opposed ring stent valleys 44A, 46A. The ring stent 22 can fold about its peaks 40A, 42A and valleys 44A, 46A to adopt a compact folded configuration, but the resilience of the ring stent urges it towards the expanded position shown in FIGS. 1 to 3. Similarly the second ring stent 24 has two diametrically opposed ring stent peaks 40B, 42B and two diametrically opposed ring stent valleys 44B, 46B, and can fold in the same way about its peaks and valleys. The folding of at least some of the peaks and valleys of the first and second ring stents 22, 24 can be selectively controlled by control wires, as is described below. The remaining stents are also capable of folding about their peaks and valleys so that they can adopt a compact folded configuration when held in a retractable sleeve 36, but their peaks and valleys are not selectively foldable.

Adjacent to the ring stent peaks 40, 42 of the first and second ring stents 22, 24 are provided hooks 48 which engage with the lumen in which the stent graft 20 is deployed when the ring stent peaks 40, 42 expand away from the delivery shaft 30. Also adjacent to the ring stent peaks 40, 42 of the first and second ring stents 22, 24 are provided connection loops 146, which serve as stent peak connection points, and through which stent peak control wires 50, 52, 54, 56 are connected, as described below. Further connection loops 148 are provided adjacent to the ring stent valleys 44, 46 of the first ring stent 22. The loops 148 serve as stent valley connection points for the connection of stent valley control wires 70, 72, as described below. Instead of providing stent valley connection points at the first ring stent 22, as illustrated in FIGS. 1 to 3, they may be provided at the second ring stent 24, or at both the first and

second ring stents **22**, **24**, or at both the first and third ring stents **22**, **24**, or on the fabric of the sleeve **28** on the valley line between the first and second ring stents **22**, **24**. The control wires may be of any suitable material and in a preferred embodiment are of Ultra High Molecular Weight Polyethylene.

The delivery apparatus **10** is shown in FIGS. **4** to **6**. The delivery apparatus **10** comprises a delivery shaft **30** having a proximal end **33** and a distal end **34**. Surrounding the delivery shaft is a retractable sleeve **36** which can be retracted using a retraction wheel **120** or a manual slider **122**, which serve to retract the retractable sleeve **36** inside the shaft **124** of the apparatus. The retraction mechanism, including the retraction wheel **120** and manual slider **122**, does not form part of the present invention, and is described in more detail in the applicant's UK patent application GB1109316.8 filed 3 Jun. 2011.

At the distal end of the shaft **124** is a grip portion **126**, which the operator holds in one hand **128** while operating the retraction wheel **120** with the other hand **129** to retract the sleeve **36**. Beyond the grip portion **126** is the control handle **38** which is used to control the deployment and position of the first and/or second ring stents **22**, **24**. The control handle **38** includes a rotary collar **58** which serves as a peak controller and two sliders **74**, **76** which serve as valley controllers.

The use of the delivery apparatus **10** to deploy a ring stent **22** of a stent graft **20** in the thoracic arch **100** will now be described with reference to FIGS. **7** to **13**, which show the thorax and apparatus schematically as well as images of the sort produced by a suitable medical imaging device. Although the invention is described with reference to the thoracic arch, it is to be understood that the apparatus and method of the invention may be used in any location in a lumen **100** where control of the angle and position of a ring stent is required, and the invention is not limited to use with the thorax.

A guide wire **32** is located to guide the delivery apparatus **10**. An intended landing zone **102** for the ring stent **22** at the end of the stent graft **20** is identified. Preferably the intended landing zone is perpendicular to the vessel wall **104**, **106** on both the inner and outer aspects of the thoracic arch. The delivery shaft **30**, on which the stent graft **20** is held by the retractable sleeve **36**, is slowly advanced along the guide wire **32** until the ring stent **22**, which is in its folded state within the retractable sleeve **36**, is at the intended landing zone, as shown in FIG. **7**.

The retraction wheel **120** is then slowly rotated, as shown in FIG. **8**, to retract the sleeve **36** to the position shown in FIG. **9**. The first ring stent **22** is prevented from unfolding by the stent peak control wires **50**, **52** which hold the ring stent peaks **40A**, **42A** towards the delivery shaft **30**. However the ring stents **26** which are not restrained by control wires are free to expand as the sleeve **36** is retracted so that the stent graft **20** adopts a tapered balloon shape. At this stage some longitudinal movement or rotation of the apparatus **10** and delivery shaft **30** is possible to ensure correct orientation and positioning of the ring stent peaks **40**, **42** and valleys **44**, **46**, since the hooks **48** which engage with the internal surface of the lumen **100** must be orientated correctly. Once the required position and orientation have been achieved, the stent graft **20** can be unsheathed from the sleeve **36** more rapidly by sliding the manual slider **122**, until the stent graft **20** is fully unsheathed as shown in FIG. **10**. The release wire **140** associated with the retractable sleeve can then be removed.

The position of the proximal ring stent **22** can now be adjusted to ensure that the ring stent **22** is positioned perpendicular to the lumen **100**. The positions of the ring stent valleys **44**, **46** are optimised by opening the valleys so that they are positioned perpendicular to the lumen wall **104**, **106**. The first ring stent valley **44** can be opened and collapsed by moving the corresponding first valley controller slider **74** forwards in the direction of arrow **134**, as shown in FIG. **11**, and backwards respectively. Similarly the second ring stent valley **46** can be opened and collapsed by moving the corresponding second valley controller slider **76** forwards in the direction of arrow **136**, as shown in FIG. **11**, and backwards respectively. Movement of the sliders **74**, **76** results in movement of the corresponding stent valley control wires **70**, **72** which extend from the control handle **38** along longitudinal passages **90** in the delivery shaft **30** and out of the delivery shaft **30** through apertures **132** to connection points **148** on the first and second ring stent valleys **44**, **46** respectively. Each valley **44**, **46** can be opened and closed independently of the other by moving the corresponding slider **74**, **76**. The resilience of the ring stent **22** urges the valleys **44**, **46** towards the open position, but the sliders **74**, **76** are provided with a ratchet **80**, seen in FIGS. **15** and **18**, so that they can only be moved when the operator releases the slider **74**, **76** from the ratchet **80** by pressing on the actuator surface **78** of the slider **74**, **76**.

Once the valley positions are optimised, the peaks **40**, **42** of the ring stent **22** can be opened so that the proximal portion of the stent graft **20** is engaged with the vessel wall **104**, **106**. The peaks **40**, **42** are opened by depressing one or more buttons or tabs **59** on the peak controller rotary collar **58** and turning it in a clockwise direction in the direction of arrow **67** about the longitudinal axis **66** of the control handle, as shown in FIG. **12**. Rotation of the collar **58** results in rotation of a peak controller winder **68** which permits movement of corresponding stent peak control wires **50**, **52** which extend from the control handle **38** along longitudinal passages **90** in the delivery shaft **30** and out of the delivery shaft **30** through apertures **130** to connection points **146** on the first and second ring stent peaks **40A**, **42A** respectively, to permit the peaks to move away from the delivery shaft **30** under the resilience of the ring stent **22**. Instead of only the peaks **40A**, **42A** of the first ring stent **22** being controllable, the peaks **40B**, **42B** of the second ring stent **24** or other ring stents **26** may also be connected by associated stent peak control wires **54**, **56** coupled in the same way to the peak controller winder **68**, so that on rotation of the collar **58**, the peaks **40B**, **42B** of the second ring stent **24** also move away from the delivery shaft **30** under the resilience of the ring stent **22**.

The resilience of the ring stents **22**, **24** urges the peaks **40**, **42** towards the open position, but the button **59** prevents rotation of the collar **58** unless the button **59** is depressed, by means of a ratchet (not shown) so that the peaks **40**, **42** can only be moved when the operator depresses the button **59**.

Once the peaks **40**, **42** are fully opened the position of the ring stent **22** and stent graft **20** can be checked. If the position needs further adjustment, the peaks **40**, **42** are retracted again to the collapsed position on the delivery shaft **30** by rotating the peak controller **58** in an anti-clockwise direction opposite to that of arrow **67**, and then the valleys **44**, **46** are similarly retracted by moving the sliders in the direction opposite to that of arrows **134**, **136**. The valleys and peaks can then be realigned to achieve a more accurate position. As before, once the valley positions are satisfactory, the peaks are opened to allow the stent graft **20** to engage with the vessel walls **104**, **106** by means of the hooks

48. The release wires **140**, whose proximal ends **96** hold the proximal end **92** of the stent graft **20** on the delivery shaft **30**, and the looped back tether wires **98**, which hold the distal end **94** of the stent graft **20** on the delivery shaft **30**, can then be removed, prior to re-sheathing with the retractable sleeve **36** and removal of the delivery apparatus from the lumen **100**. The proximal end **96** of each release wire **140** is associated with a connection **146, 148** between the stent graft **20** and one of the stent peak control wires **50, 52**, the stent valley control wires **70, 72**, and the tow or back tether wires **98**, so that pulling the release wire **140** causes the connection **146, 148** to be broken. Such release wires **140** are known, for example in EP-A-910309, and are not described further. In the described embodiment the release wires travel through the proximal end **33** of delivery shaft **30** to assist in providing stability during the removal of the release wires. The distal ends **144** of the release wires **140** may each be secured to a release clip **154** mounted at the distal end of the control handle **38**. The release wires **140** associated with the stent peak control wires **50, 52** and the stent valley control wires **70, 72** are also removed by pulling the corresponding release clip **154** distally along the longitudinal axis **66** of the control handle, as shown in FIG. **14**.

The interior of the control handle is shown more clearly in FIGS. **15** to **18** with the handle access cover removed and with the control wires omitted for clarity. The control wires **50, 52, 54, 56, 70, 72** pass from the proximal end **33** of the delivery shaft **30** through one or more longitudinal passages in the delivery shaft **30** to the distal end **34** where they enter the shaft **124** of the control handle **38** through a central passage **127**. The control wires **50, 52, 54, 56, 70, 72** then pass through a manifold **60** having a discrete tube aperture **62** for each control wire. This provides a smooth and continuous passage for each control wire from the connection point **146, 148** at the stent graft **20** to the peak controller **58** or valley controllers **74, 76**. The manifold **60** enables feeder wires (not shown) to pass freely for loading the control wires when assembling the apparatus. Integral haemostasis valves are provided in a haemostasis switching plate **64** adjacent to the manifold **60** to allow flushing of the system with saline prior to use. The haemostasis valves also provide a controlled restriction to prevent excessive blood loss while maintaining control of the stent graft **20** by the apparatus. The manifold apertures may comprise passages which diverge from the connection with central passage **127** to the distal side of the haemostasis switching plate **64**, to enable the control wires to be distributed and loaded onto the peak controller **58** or valley controllers **74, 76**.

The valley controllers **74, 76** each include an actuating surface **78** which must be depressed to disengage the slider **74, 76** from the ratchet **80** to allow the slider to be moved. Once pressure is removed from the actuating surface **78**, the slider **74, 76** becomes engaged again with the ratchet **80** and further movement of the slider **74, 76** is prevented. The valley controllers **74, 76** each include a connector **82** to which the corresponding stent valley control wire **70, 72** is connected.

The peak controller **58** includes a winder **68** (omitted in FIG. **15** but shown in FIGS. **16** to **20**) which is mounted for rotation on a winder spindle **69**. The rotary collar **58** is omitted for clarity in FIGS. **15** to **20**. A ratchet plate (not shown) is mounted on the winder spindle **69** and provides bearing and ratchet interfaces to the winder **68**.

The connection of the control wires **50, 52, 54, 56, 70, 72** is shown in FIGS. **19** to **22**. Each wire may comprise a looped wire, having two strands, but is referred to in this specification as a single wire. At the distal end the stent

valley control wires **70, 72** pass out of the manifold apertures **62** in the control handle **38** and are connected to the slider wire connectors **82**. At the proximal end the stent valley control wires **70, 72** pass out of the valley apertures **132** in the delivery shaft **38** and are connected to the connection points **148** on the ring stent valleys **44, 46**. In the illustrated example of FIG. **22** the stent valley control wires **70, 72** are connected to the connection points **148** on the valleys **44B, 46B** on the second ring stent **24**, but they could instead be connected to the valleys **44A, 46A** on the first or third ring stent **22, 24**, or to the valleys of two ring stents **22, 24, 26**, or to the sleeve **28** between the ring stents **22, 24, 26**.

At the distal end the stent peak control wires **50, 52, 54, 56** pass out of the manifold apertures **62** in the control handle **38**, pass through apertures **160** and are connected to connection points on the peak controller winder **68**. In the illustrated example the connection points are anchor pins or fixing clamps **162** which enable adjustment of the control wires to achieve the correct length. At the proximal end the stent peak control wires **50, 52, 54, 56** pass out of the peak apertures **130** in the delivery shaft **38** and are connected to the connection points **146** on the ring stent peaks **40A, 42A, 40B, 42B**. In the illustrated example the stent peak control wires are connected to the connection points **146** on the peaks of the both the first and second ring stents **22, 24**, but they could instead be connected to the peaks on the first ring stent **22** only, or the peaks of the second ring stent **24** only.

The apparatus and method of the invention allow the controlled rotation and orientation of a ring stent **22, 24** in a lumen **100** by selectively operating one of the first and second valley controllers **74, 76** to permit movement of the stent valley control wires **70, 72** along the delivery shaft **30** such that the first or second diametrically opposed valley connection point **148** on the ring stent **22, 24** is urged by the resilience of the ring stent to move away from the delivery shaft **30**, thereby rotating the ring stent. The movement is reversible so that the valley connection point **148** can also be moved towards the delivery shaft **30** to a folded or contracted position. During assembly of the apparatus the stent graft **20** adopts its expanded position shown in FIG. **22**, but prior to sheathing the two stent valley control wires **70, 72** are tensioned such that the valley connection points **148** are held by the stent valley control wires **70, 72** in a folded arrangement towards the delivery shaft. In addition the two stent peak control wires **50, 52, 54, 56** are tensioned such that the peak connection points **146** are also held in a folded arrangement towards the delivery shaft.

The apparatus and method of the invention allow the controlled deployment of a ring stent **22, 24** in a lumen **100** by operating the peak controller **58** to permit movement of the stent peak control wires **50, 52, 54, 56** along the delivery shaft such that the peak connection points **146** on one or more proximal ring stents **22, 24** are urged by the resilience of the ring stent to move away from the delivery shaft and the ring stent adopts a fully deployed position. The movement is reversible so that the peak connection point **146** can also be moved towards the delivery shaft **30** to a folded or contracted position if the ring stent requires repositioning.

The peak and valley connection points **146, 148** may be eyelets through which the looped control wires are threaded.

All documents referred to in this specification are herein incorporated by reference. Various modifications and variations to the described embodiments of the inventions will be apparent to those skilled in the art without departing from the scope and spirit of the invention. Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as

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claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes of carrying out the invention which are obvious to those skilled in the art are intended to be covered by the present invention.

REFERENCE NUMERALS

10 delivery apparatus
 20 tubular prosthesis (stent graft)
 22 ring stent
 24 second ring stent
 26 other ring stents
 28 fabric sleeve
 30 delivery shaft
 31 central passage
 32 guide wire
 33 proximal end
 34 distal end
 36 retractable sleeve
 38 control handle
 40A, 40B ring stent peak
 42A, 42B ring stent peak
 44A, 44B ring stent valley
 46A, 46B ring stent valley
 48 hooks
 50, 52, 54, 56 stent peak control wire
 58 peak controller (rotary collar)
 59 button on collar
 60 control handle manifold
 62 control handle manifold apertures
 64 haemostasis switching plate with manifold haemostasis valves
 66 longitudinal axis of the control handle
 67 arrow
 68 peak controller winder
 69 winder spindle
 70, 72 stent valley control wire
 74, 76 valley controller (slider)
 78 valley controller actuating surface
 80 valley controller ratchet
 82 slider wire connector
 90 delivery shaft longitudinal passages
 92 proximal end of a tubular prosthesis
 94 distal end of a tubular prosthesis
 96 release wire proximal ends
 98 back tether wires
 100 lumen
 102 intended location
 104, 106 inner surface of lumen
 120 retraction wheel
 122 manual slider
 124 apparatus shaft
 126 grip portion
 127 central passage
 128, 129 hands
 130 delivery shaft peak apertures
 132 delivery shaft valley apertures
 134, 136 arrow
 140 release wires
 144 release wire distal ends
 146, 148 connection between a stent graft and stent peak control wire or stent valley control wire
 154 release wire release clip
 160 aperture
 162 fixing clamp

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The invention claimed is:

1. A delivery apparatus, the apparatus comprising:
 - a delivery shaft with a tubular prosthesis mounted on a proximal end thereof for insertion into a lumen and a distal end remote from the proximal end, the prosthesis comprising a sleeve and at least one discrete saddle-shaped ring stent with opposing peaks and opposing valleys positioned between the peaks, the at least one discrete saddle-shaped ring stent is attached to the sleeve to maintain the patency of the sleeve after deployment of the prosthesis;
 - a retractable sleeve adapted to contain the tubular prosthesis mounted on the delivery shaft;
 - a control handle adjacent the distal end of the delivery shaft;
 - at least one stent peak control wire extending from the control handle and directly attached to at least one peak of the opposing peaks at an at least one peak connection point;
 - at least one peak controller at the control handle adapted to control the position of the at least one stent peak control wire to selectively open the at least one peak from a collapsed state and collapse the at least one peak from an open state;
 - at least one stent valley control wire extending from the control handle and directly attached to at least one valley of the opposing valleys at an at least one valley connection point; and
 - at least one valley controller at the control handle adapted to control the position of the at least one stent valley control wire to selectively open the at least one valley from a collapsed state and collapse the at least one valley from an open state.
2. The apparatus of claim 1, comprising two or more stent peak control wires extending from the control handle, and wherein the at least one peak controller is adapted to control the position of the two or more stent peak control wires.
3. The apparatus of claim 1, comprising two stent valley control wires extending from the control handle, and wherein the apparatus includes two valley controllers at the control handle, each valley controller being adapted to control the position of a respective stent valley control wire of the two stent valley control wires.
4. The apparatus of claim 1, wherein the delivery shaft includes a plurality of longitudinal passages, each stent peak control wire and stent valley control wire extending through a respective longitudinal passage.
5. The apparatus of claim 1, including one or more tow wires extending from the proximal end of the delivery shaft to a proximal end of the tubular prosthesis mounted on the delivery shaft.
6. The apparatus of claim 1, including one or more back tether wires extending from the proximal end of the delivery shaft to a distal end of the tubular prosthesis mounted on the delivery shaft.
7. The apparatus of claim 1, wherein the control handle includes a manifold having a plurality of apertures, each stent peak control wire and stent valley control wire extending through a respective aperture.
8. The apparatus of claim 7, wherein the manifold includes a plurality of haemostasis valves switchable between open and closed positions.
9. The apparatus of claim 1, wherein the at least one peak controller comprises a rotatable collar.
10. The apparatus of claim 9, wherein the rotatable collar is mounted for rotation about a longitudinal axis of the control handle.

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11. The apparatus of claim 9, wherein the at least one peak controller includes a winder which rotates with the collar, and the at least one stent peak control wire being wound on the winder.

12. The apparatus of claim 9, wherein rotation of the rotatable collar in a first direction opens the at least one ring stent, and rotation of the rotatable collar in a second direction collapses the at least one ring stent.

13. The apparatus of claim 1, wherein the at least one valley controller comprises a slider.

14. The apparatus of claim 13, wherein the at least one valley controller includes an actuating surface and a ratchet adapted to allow the slider to move when the actuating surface is depressed and to hold the slider in an engaged position when the actuating surface is released.

15. The apparatus of claim 13, wherein the slider is mounted for longitudinal sliding movement.

16. The apparatus of claim 13, wherein the slider includes a wire connector to which is connected the at least one stent valley control wire.

17. The apparatus of claim 13, wherein sliding movement of the slider in a first direction opens the at least one ring stent, and sliding movement of the slider in a second direction collapses the at least one ring stent.

18. The apparatus of claim 1, wherein the delivery shaft includes a plurality of peak apertures adjacent the proximal end thereof, the at least one stent peak control wire extending through one of the plurality of peak apertures.

19. The apparatus of claim 1, wherein the delivery shaft includes a plurality of valley apertures adjacent the proximal end thereof, the at least one stent valley control wire extending through one of the plurality of valley apertures.

20. The apparatus of claim 19, wherein the delivery shaft further includes a plurality of peak apertures, and wherein the valley apertures are arranged between the peak apertures and the distal end.

21. The apparatus of claim 1, wherein the at least one stent valley control wire and the at least one stent peak control wire each comprise at least one looped wire which extends from the control handle to the proximal end of the delivery shaft and back to the control handle.

22. The apparatus of claim 1, further including a plurality of release wires extending from the control handle along the delivery shaft and having a proximal end at or near the proximal end of the delivery shaft and a distal end at the control handle.

23. The apparatus of claim 22, wherein the distal end of each release wire is connected to a release clip mounted on the control handle.

24. The apparatus of claim 23, wherein the proximal end of each release wire is connected to the at least one peak connection point or the at least one valley connection point.

25. The apparatus of claim 24, wherein the release clip is removable from the control handle to release the respective at least one peak or valley connection point.

26. The apparatus of claim 1, wherein the distance between the at least one valley connection point and the control handle is less than the distance between the at least one peak connection point and the control handle.

27. The apparatus of claim 1, wherein at least one of:
the at least one peak connection point comprises at least one point on the at least one peak of the at least one ring stent; and
the at least one valley connection point comprises at least one point on the at least one valley of the at least one ring stent.

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28. The apparatus of claim 1, wherein the at least one peak connection point comprises at least one point on the at least one peak of the at least one ring stent.

29. The apparatus of claim 28, wherein the at least one valley connection point comprises at least one point on the at least one valley of the at least one ring stent.

30. The apparatus of claim 1, wherein the selectively opening and collapsing the at least one peak comprises selectively opening and collapsing of a portion of the at least one ring stent, and the selectively opening and collapsing the at least one valley comprises selectively opening and collapsing a portion of the at least one ring stent.

31. The apparatus of claim 1, wherein the at least one ring stent of the tubular prosthesis is mounted on the delivery shaft in a collapsed configuration and is naturally biased into an open configuration from the collapsed configuration.

32. The apparatus of claim 1, wherein the sleeve is a flexible tubular conduit.

33. The apparatus of claim 32, wherein the sleeve is formed of woven or knitted flexible fabric.

34. The apparatus of claim 32, wherein the prosthesis comprises a plurality of the discrete saddle-shaped ring stents attached to the sleeve at spaced intervals.

35. The apparatus of claim 1, wherein at least one of:
the at least one peak connection point comprises at least one peak connection loop; and
the at least one valley connection point comprises at least one valley connection loop.

36. The apparatus of claim 1, wherein at least one of:
the at least one peak connection point comprises at least one peak eyelet, and the at least one stent peak control wire is threaded through the at least one peak eyelet; and
the at least one valley connection point comprises at least one valley eyelet, and the at least one stent valley control wire is threaded through the at least one valley eyelet.

37. The apparatus of claim 1, wherein the at least one stent peak control wire comprises a first stent peak control wire and a second stent peak control wire, and the at least one stent valley control wire comprises a first stent valley control wire and a second stent valley control wire, wherein the at least one peak connection point comprises a first peak connection point and a second peak connection point, and the at least one valley connection point comprises a first valley connection point and a second valley connection point, wherein the opposing peaks comprise a first peak and a second peak, and the opposing valleys comprise a first valley and a second valley, wherein the first stent peak control wire is directly attached to the first peak at the first peak connection point, and the second stent peak control wire is directly attached to the second peak at the second peak connection point, and wherein the first stent valley control wire is directly attached to the first valley at the first valley connection point, and the second stent valley control wire is directly attached to the second valley at the second valley connection point.

38. The apparatus of claim 37, wherein the at least one valley controller comprises a first valley controller and a second valley controller, and wherein the first valley controller is adapted to control the position of the first stent valley control wire to selectively open the first valley from a collapsed state and collapse the first valley from an open state, and the second valley controller is adapted to control the position of the second stent valley control wire to selectively open the second valley from a collapsed state and collapse the second valley from an open state.

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39. A method, comprising:
 providing the delivery apparatus of claim 1;
 holding the at least one stent peak control wire under
 tension such that the at least one peak connection point
 is held by the at least one stent peak control wire and
 the at least one peak is held in a folded arrangement
 towards the delivery shaft; and
 selectively operating the at least one peak controller to
 permit movement of the at least one stent peak control
 wire along the delivery shaft such that the at least one
 peak is urged by resilience of the at least one ring stent
 to move away from the delivery shaft and the at least
 one ring stent adopts a partially deployed position.
40. The method of claim 39, further comprising:
 holding the at least one stent valley control wire under
 tension such that the at least one valley connection
 point is held by the at least one stent valley control wire
 and the at least one valley is held in a folded arrange-
 ment towards the delivery shaft; and
 selectively operating the at least one valley controller to
 permit movement of the at least one stent valley control
 wire along the delivery shaft such that the at least one
 valley is urged by the resilience of the at least one ring
 stent to move away from the delivery shaft and to rotate
 the at least one ring stent in the partially deployed
 position.
41. The method of claim 40, wherein the selectively
 operating the at least one valley controller comprises mov-
 ing a slider which includes a wire connector to which is
 connected the at least one stent valley control wire.
42. The method of claim 40, wherein the at least one
 valley controller comprises a second valley controller, the at
 least one stent valley control wire comprises a second stent
 valley control wire, and the at least one valley comprises a
 second valley, and wherein the method further comprises
 selectively operating the second valley controller to permit
 movement of the second stent valley control wire along the
 delivery shaft such that the second valley is urged by the
 resilience of the at least one ring stent to move away from
 the delivery shaft and the at least one ring stent adopts a
 deployed or selected rotational position.
43. The method of claim 39, wherein the at least one
 valley connection point comprises at least one point on the
 at least one valley of the at least one ring stent.
44. The method of claim 39, wherein at least one of the
 at least one peak connection point and the at least one valley
 connection point comprises at least one eyelet.
45. The method of claim 39, wherein the selectively
 operating the at least one peak controller includes rotating a
 rotatable collar to wind or unwind the at least one stent peak
 control wire on a winder.
46. The method of claim 39, wherein the at least one stent
 valley control wire and the at least one stent peak control
 wire each comprise at least one looped wire which extends
 from the control handle to the at least one valley connection
 point and the at least one peak connection point, respec-
 tively, and back to the control handle.
47. The method of claim 39, wherein at least one of:
 the at least one peak connection point comprises at least
 one peak connection loop; and
 the at least one valley connection point comprises at least
 one valley connection loop.
48. The method of claim 39, wherein at least one of:
 the at least one peak connection point comprises at least
 one peak eyelet, and the at least one stent peak control
 wire is threaded through the at least one peak eyelet;
 and

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- the at least one valley connection point comprises at least
 one valley eyelet, and the at least one stent valley
 control wire is threaded through the at least one valley
 eyelet.
49. The method of claim 48, wherein at least one of the
 at least one stent peak control wire and the at least one stent
 valley control wire comprises a looped wire which extends
 through the at least one eyelet of the at least one peak
 connection point or the at least one valley connection point,
 respectively, and back to the control handle.
50. A method, comprising:
 providing the delivery apparatus of claim 1;
 holding the at least one stent two valley control wire under
 tension such that the at least one valley connection
 point is held by the at least one stent valley control wire
 and the at least one valley is held in a folded arrange-
 ment towards the delivery shaft; and
 selectively operating the at least one valley controller to
 permit movement of the at least one stent valley control
 wire along the delivery shaft such that the at least one
 valley is urged by resilience of the at least one ring stent
 to move away from the delivery shaft and to rotate the
 at least one ring stent.
51. The method of claim 50, wherein at least one of:
 the at least one peak connection point comprises at least
 one point on the at least one peak of the at least one ring
 stent; and
 the at least one valley connection point comprises at least
 one point on the at least one valley of the at least one
 ring stent.
52. The method of claim 50, further comprising:
 holding the at least one stent peak control wire under
 tension such that the at least one peak connection point
 is held by the at least one stent peak control wire and
 the at least one peak is held in a folded arrangement
 towards the delivery shaft; and
 selectively operating the at least one peak controller to
 permit movement of the at least one stent peak control
 wire along the delivery shaft such that the at least one
 peak is urged by the resilience of the at least one ring
 stent to move away from the delivery shaft and the at
 least one ring stent adopts a partially deployed position.
53. The method of claim 50, wherein at least one of the
 at least one peak connection point and the at least one valley
 connection point comprises at least one eyelet.
54. The method of claim 50, wherein the selectively
 operating the at least one valley controller comprises mov-
 ing a slider which includes a wire connector to which is
 connected the at least one stent valley control wire.
55. The method of claim 50, wherein the at least one
 valley controller comprises a second valley controller, the at
 least one stent valley control wire comprises a second stent
 valley control wire, and the at least one valley comprises a
 second valley, and wherein the method further comprises
 selectively operating the second valley controller to permit
 movement of the second stent valley control wire along the
 delivery shaft such that the and second valley is urged by the
 resilience of the at least one ring stent to move away from
 the delivery shaft and the at least one ring stent adopts a
 deployed or selected rotational position.
56. The method of claim 50, wherein at least one of:
 the at least one peak connection point comprises at least
 one peak connection loop; and
 the at least one valley connection point comprises at least
 one valley connection loop.

57. The method of claim 50, wherein at least one of:
the at least one peak connection point comprises at least
one peak eyelet, and the at least one stent peak control
wire is threaded through the at least one peak eyelet;
and
the at least one valley connection point comprises at least
one valley eyelet, and the at least one stent valley
control wire is threaded through the at least one valley
eyelet.

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58. The method of claim 57, wherein the at least one stent
valley control wire comprises at least one looped wire which
extends through the at least one eyelet and back to the
control handle.

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UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

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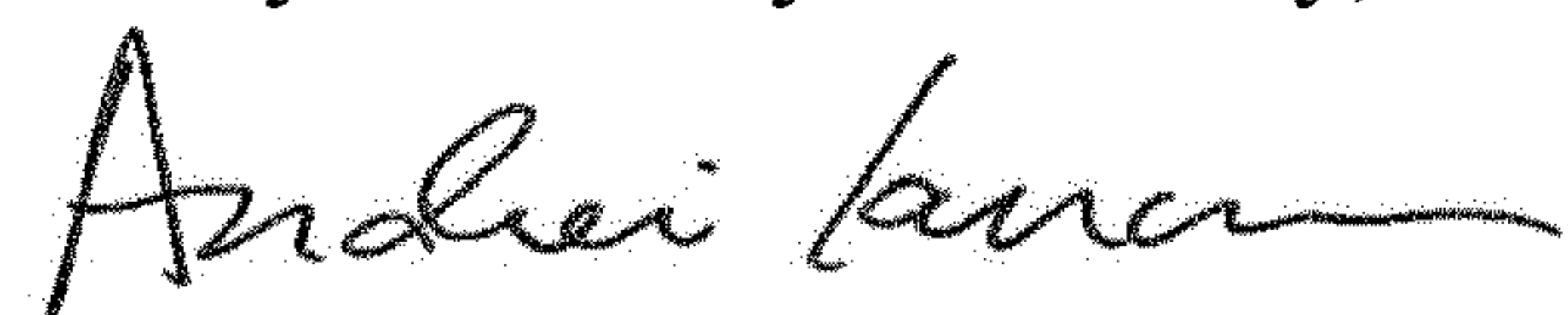
Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In the Claims

Column 18, Line 13: Claim 50, Delete "one stent two valley" and insert -- one stent valley --

Signed and Sealed this
Twenty-ninth Day of January, 2019



Andrei Iancu
Director of the United States Patent and Trademark Office